

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:20:48 ; Search time 307.385 Seconds

(without alignments)
10509.347 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atggttgatcttcttgcctt.....gcctgagtgctgcttact 111Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

```
GenEmbl:*
1: gb_ba:*
2: gb_htg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pin:*
35: em_htg_rnd:*
36: em_htg_mam:*
37: em_htg_vtl:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*
```

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	111	100.0	169620	2	AC012674	AC012674 Homo sapi
2	48	43.2	143372	9	AL137847	AL137847 Homo DNA
3	40	36.0	123779	30	AC021025	Ac021025 Homo sapi
4	40	36.0	128118	2	AC076969	AC076969 Homo sapi
5	40	36.0	128583	9	AC121249	AC121249 Homo sapi
6	40	36.0	148290	9	AC107979	AC107979 Homo sapi
7	40	36.0	165649	9	AC103996	AC103996 Homo sapi
8	40	36.0	178650	9	AC104303	AC104303 Homo sapi
9	40	36.0	192826	9	AC080762	AC080762 Homo sapi
10	39	35.1	32918	2	AC007445	AC007445 Homo sapi
11	39	35.1	38936	9	AL358817	AL358817 Human DNA
12	39	35.1	124271	2	AC025179	AC025179 Homo sapi
13	39	35.1	146671	9	AC008814	AC008814 Homo sapi
14	39	35.1	159747	2	AP001019	AP001019 Homo sapi
15	39	35.1	162740	2	AC034249	AC034249 Homo sapi
16	39	35.1	169772	9	AC069538	AC069538 Homo sapi
17	39	35.1	175466	9	AL607077	AL607077 Human DNA
18	38	34.2	110000	2	AL691517_2	Continuation (3 of
19	38	34.2	166706	9	AC068875	AC068875 Homo sapi
20	38	34.2	207408	2	AC087283	AC087283 Homo sapi
21	38	34.2	207548	9	AL591491	AL591491 Human DNA
22	36	32.4	121720	9	AL591491	AL591491 Human DNA
23	35	31.5	153940	9	AC022294	AC022294 Homo sapi
24	35	31.5	325069	2	AC079737	AC079737 Homo sapi
25	34	30.6	99577	9	AC026324	AC026324 Homo sapi
26	34	30.6	108040	2	AC068150	AC068150 Homo sapi
27	34	30.6	134760	9	AC099484	AC099484 Homo sapi
28	34	30.6	146059	9	AC019030	AC019030 Homo sapi
29	34	30.6	172206	9	AC092119	AC092119 Homo sapi
30	34	30.6	172567	2	AC015493	AC015493 Homo sapi
31	34	30.6	173166	9	AC092375	AC092375 Homo sapi
32	34	30.6	273807	2	AC025421	AC025421 Homo sapi
33	34	30.6	316296	2	AC092285	AC092285 Homo sapi
34	33	29.7	33458	9	HS2160101	AL109656 Human DNA
35	33	29.7	49616	9	AL365267	AL365267 Human DNA
36	33	29.7	77743	9	HSTCRBV	U03115 Human V bet
37	33	29.7	153788	2	AC023971	AC023971 Homo sapi
38	33	29.7	166434	9	AC020717	AC020717 Homo sapi
39	33	29.7	215422	9	U66060	U66060 Human germ
40	31	27.9	70313	2	AC016216	AC016216 Homo sapi
41	31	27.9	82225	9	HS253014	280771 Human DNA
42	31	27.9	164550	2	CNS01RHY	AL162633 Human chr
43	31	27.9	173706	2	AC022694	AC022694 Homo sapi
44	31	27.9	176418	9	AC090811	AC090811 Homo sapi
45	31	27.9	184759	9	AC022695	AC022695 Homo sapi

ALIGNMENTS

```
RESULT 1
LOCUS      AC012674/c
DEFINITION Homo sapiens chromosome 3 clone RPI-458H3, WORKING DRAFT SEQUENCE,
ACCESSION  AC012674
VERSION    AC012674.10 GI:9719580
KEYWORDS   HTG: HTGS_PHASE1, HTGS_DRAFT.
SOURCE     Homo sapiens.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 169620)
            Muzny,D.M., Adams,C., Bailey,M., Barbara,J., Blankenburg,K.,
            Bodola,B., Bouck,J., Bowie,S., Brooks,A., Buhay,C., Bunac,C.,
```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N., Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D., Forcum-Tansey, J., Frantz, P., Ganesh, R., Gorrell, J.H., Gorrell, L.L., Goulet, W., Harris, K., Hernandez, J., Hodgson, A., Hughes, M., Hollaway, C., Hosak, H., Jackson, L.E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondejowski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Licharge, O., Liu, J., Liu, W., Logan, O., Lozano, R.J., Lu, J., Lucier, R., Martin, R., Martinez, C., McLeod, M.P., Mei, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Oswald, G., Parish, B., Paxton, S., Payton, B., Perez, L., Pu, L.L., Quinn, M., Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S., Shah, E., Shen, H., Simon, M., Sparks, A., Stamps, A., Suganag, R., Taber, P., Taylor, T., Vasquez, L., Vinson, R., Vo, O., Webb, M., Watlington, S., Weinstein, G., Weinstein, I.R., Williamson, A., Worley, K., Wren, J., Wrensford, G., Yu, W., Zhou, X., Nelson, D., and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.

Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
Project Information
Center project name: HMO3
Center clone name: RPI-458H3

Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 163720; sum-of-contigs estimation
Estimated insert size: 171608; agarose-fp estimation
Quality coverage: 3.9x in Q20 bases; agarose-fp estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank/draft_data.html).
NOTE: This is a 'working draft' sequence. It currently consists of 18 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1
28689: contig of 28689 bp in length
28789: gap of unknown length
28790: contig of 22043 bp in length
50833: gap of unknown length
50933: gap of unknown length
69144: contig of 18212 bp in length
69244: gap of unknown length
69245: contig of 14960 bp in length
84204: gap of unknown length
84305: contig of 10363 bp in length
94667: gap of unknown length
94767: gap of unknown length
107261: contig of 12494 bp in length
107262: gap of unknown length
107361: gap of unknown length
107362: contig of 10189 bp in length
117550: gap of unknown length
117551: gap of unknown length
117650: contig of 9289 bp in length
126939: gap of unknown length
127038: contig of 8001 bp in length
127040: gap of unknown length
135040: contig of 8001 bp in length
135041: gap of unknown length
135140: contig of 6499 bp in length
135141

141640 141739: gap of unknown length
141740 149558: contig of 7819 bp in length
149559 149559: gap of unknown length
149562 149562: contig of 4904 bp in length
154563 154563: gap of unknown length
154662 154662: contig of 4325 bp in length
158988 159087: gap of unknown length
159088 162376: contig of 3289 bp in length
162377 162476: gap of unknown length
162477 165191: contig of 2715 bp in length
165192 165291: gap of unknown length
165292 167173: contig of 1882 bp in length
167174 167273: gap of unknown length
167274 168393: contig of 1120 bp in length
168394 168493: gap of unknown length
168494 169620: contig of 1127 bp in length.

FEATURES
source
1. 169620
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RPI-458H3"

BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others
ORIGIN

Query Match 100.0%; Score 111; DB 2; Length 169620;
Best Local Similarity 100.0%; Pred No. 2.3e-53;
Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGGGTGATCTTTTGGCTTCAGAGATCTTTTCATCTTTCAGAGACTTTCGGCCG 60
|||||
Db 87441 ATGGGTGATCTTTTGGCTTCAGAGATCTTTTCATCTTTCAGAGACTTTCGGCCG 87382
|||||

OY 61 GAGATGTAACCTCGGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 111
|||||
Db 87381 GAGATGTAACCTCGGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 87331
|||||

RESULT 2
AL137847/c
LOCUS
DEFINITION Human DNA sequence from clone RPI1-439K3 on chromosome 9q42.2-31.1.
ACCESSION AL137847
VERSION AL137847
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 143372)
Kimberley, A.
Direct Submission
Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk
On Nov 17, 2001 this sequence version replaced gi:16408610.
During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP

COMMENT

Matches 40: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

QY 69 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 108
Db 3083 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 3044

RESULT 4

AC076969

LOCUS

DEFINITION

AC076969 128118 bp DNA linear HNG 15-OCT-2001
Homo sapiens chromosome 3 clone RP11-79K12, WORKING DRAFT SEQUENCE,
14 unordered pieces.

ACCESSION

AC076969

VERSION

AC076969.6

KEYWORDS

HTG: HTGS_PHASE1, HTGS_DRAFT.

SOURCE

Homo sapiens

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 128118)

Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,

Alstrooms,S.L., Amaralunge,H.C., Are,J.R., Banks,T., Barabara,J.,

Benton,J., Bimaga,K., Blankenburg,K., Bonnin,D., Bouck,J.,

Bowle,S., Bivava,M., Brown,E., Brown,M., Bryant,N.P., Bunay,C.,

Burke,P., Burrell,C., Burrell,K.L., Byrd,N.C., Caron,T.F.,

Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,

Chen,Z., Chowdhury,I., Christopoulos,C., Cleveland,C.D., Cox,C.,

Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,

Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,

Dem,A.L., Ding,Y., Dinh,H.H., Doultwalte,K.J., Drepper,H.,

Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,

Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,

Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,

Gara-N., Gill,R., Gorrell,J.H., Guevara,M., Gunaratne,P., Hale,S.,

Hamilton,K., Harris,C., Harris,K., Hart,M., Haylak,P., Hayes,A.,

Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C.,

Hollins,B., Homsi,F., Howard,S., Huber,J., Huliy,S., Hume,J.,

Jackson,L.E., Jacobson,B., Jia,T., Johnson,R., Jollive,S.,

Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,T.,

Kovar,C., Kralovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,

Lewis,L., Li,J., Li,Z., Licharge,O., Lieu,C., Liu,J., Liu,W.,

Lousaged,H., Lozano,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,

Ma,J., Maheshwari,M., Mapa,P., Martin,R., Martindale,A.,

Martinez,E., Massey,E., Mawhney,E., McLeod,M.P., Meador,M.,

Mei,G., Metzger,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K.,

Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,N.,

Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokweto,S.,

Ogun,M., Okunolu,G., Otagunye,N., Oviedo,R., Pace,A., Payton,B.,

Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L.,

Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojebokan,I., Rolfe,M.,

Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shoostari,N.,

Slison,I., Sodergren,E., Sonake,T., Sparks,A., Stanley,H.,

Stone,H., Sutton,A., Swatek,A., Taber,P., Tamerisa,A., Tamerisa,K.,

Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N.,

Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalob,D., Vinson,R.,

Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,

Watlington,S., Williams,G., Williamson,A., Wiczysk,R., Wooden,S.,

Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,

Weinstock,G. and Gibbs,R.

BASE COUNT 35433 a 25053 c 23792 g 42367 t 1473 others
ORIGIN
Query Match 36.0%; Score 40; DB 2; Length 128118;
Best Local Similarity 100.0%; Pred. No. 4.3e-12;
Matches 40: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

QY 69 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 108
Db 55106 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 55145

RESULT 5
AC121249/c 128583 bp DNA linear PRI 01-JUN-2002
LOCUS Homo sapiens chromosome 3 clone RP11-79K17, complete sequence.
DEFINITION

----- Project Information
Center project name: HBRU
Center clone name: RP11-79K12
----- Summary Statistics
Sequencing vector: M13: 108821
Assembly program: Phrap; version 0.990329
Consensus quality: 117905 bases at least Q40
Consensus quality: 121494 bases at least Q20
Estimated insert size: 122854; sum-of-coverage estimation
Quality coverage: 0x in Q20 bases; agarose-IP estimation
Quality coverage: 4x in Q20 bases; sum-of-coverage estimation

* NOTE: Estimated insert size may differ from sequence length.
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a "working draft" sequence. It currently
* consists of 14 contigs. The true order of the pieces is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 20354: contig of 20354 bp in length
20355 20454: gap of unknown length
20455 33071: contig of 12617 bp in length
33072 33171: gap of unknown length
33172 46935: contig of 13764 bp in length
46936 47035: gap of unknown length
47036 57066: contig of 10031 bp in length
57067 70378: contig of 13212 bp in length
70379 70479: gap of unknown length
70479 81786: contig of 11308 bp in length
81787 81886: gap of unknown length
81887 91428: contig of 9542 bp in length
91429 91528: gap of unknown length
91528 100455: contig of 8927 bp in length
100456 100555: gap of unknown length
100556 107879: contig of 7324 bp in length
107880 107979: gap of unknown length
107980 114149: contig of 6170 bp in length
114150 114249: gap of unknown length
114250 120186: contig of 5937 bp in length
120187 120286: gap of unknown length
120287 123336: contig of 2950 bp in length
123337 123337: gap of unknown length
123337 125606: contig of 2270 bp in length
125607 125706: gap of unknown length
125707 128118: contig of 2412 bp in length.

FEATURES
Source 1.128118
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP11-79K12"

ACCESSION AC121249 AC021025
VERSION AC121249.1 GI:20806313
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 128583)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.,
and Haugen, E.D.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 128583)
AUTHORS Kaul, R.K., Olson, M.V., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (16-MAY-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (bases 1 to 128583)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.,
and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (01-JUN-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
On May 16, 2002 this sequence version replaced gi:9719675.
----- Genome Center
Center: University of Washington Genome Center
Center Code: UMG
Web site: http://www.genome.washington.edu
Contact: uwgchgs@u.washington.edu
Drafting Center: BCM
----- Project Information
Center project name: chr-3
Center clone name: RP11-79K17 (bc0196)
----- Summary Statistics: 45% of reads
Sequencing vector: Plasmid; 55% of reads
Sequencing vector: M13; L08821; 55% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 128548 bases at least Q40
Consensus quality: 128580 bases at least Q30
Consensus quality: 128581 bases at least Q20
Insert size: 128583; sum-of-contigs
Quality coverage: 13.4x in Q20 bases; sum-of-contigs

Overlapping Sequences:
5' RP11-147N17 (UMGC:bc0267) AC104300, 22116-bp overlap
3' RP11-391P4 (UMGC:bc0402) AC104303, 114502-bp overlap

Sequence Quality Assessment:
This entry has been annotated with sequence quality
estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero.
Quality levels above 40 are expected to have less than
1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
Genbank flat file format but are available as part
of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:
all regions were either double-stranded or sequenced with an
alternate chemistry or covered by high quality data (i.e., Phred
quality >= 30); an attempt was made to resolve all sequencing
problems, such as compressions and repeats; all regions were
covered by at least one plasmid subclone or more than one M13
subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:
This sequence has been validated by Multiple Complete Digest
fingerprinting. Comparison of the experimentally derived digest
fragments with sequence-predicted fragments is given below.
The electronically-digested sequence consists of both insert and

vector, in order to accurately represent the entire circular BAC.
Small fragments below a variable cutoff (approximately 400-800 bp)
are not resolved in the fingerprint and hence do not appear
in the table. There are no significant remaining discrepancies
between the experimental and predicted values. Uniquely ordered
fragments are separated by dashed lines.

NsII		BgIII		EcoRI	
SeqDerMap	FingerPrint	SeqDerMap	FingerPrint	SeqDerMap	FingerPrint
12049	12136	4077	4013	8696	8902
579	<800	2067	2145	6	<800
1056	1065	9698	9932	1323	1337
13085	13126	3729	3734	1024	961
10944	10836	1632	1616	8421	8408
581	<800	8069	8139	1114	1126
882	887	1507	1485	3538	3534
8435	8467	11939	11871	6051	6022
3924	4103	133	<800	5129	5048
9842	9759	2135	2145	58	<800
4401	4377	78	<800	1895	1903
4458	4377	5873	5861	799	<800
4587	4522	896	902	858	885
6048	5947	3312	3356	931	885
1395	1385	1505	1485	859	885
3059	3071	6694	6716	3934	3936
3911	3861	10032	9932	1065	1058
138	<800	2168	2145	2629	2723
50	<800	1794	1785	859	885
582	<800	227	<800	6855	6888
2225	2229	4002	4013	7821	7819
984	967	2714	2726	7210	7201
887	887	5628	5624	6992	6888
1042	1065	743	758	1204	1224
2783	2808	3045	2963	2722	2723
7074	7158	1288	1262	321	<800
4087	3861	967	973	1066	1058
7976	7926	461	<800	470	<800
1233	1199	4649	4618	3055	3072
3875	3861	5289	5417	10961	10848
27	<800	1098	1095	6725	6888

1137	1065	798	809	1896	1903
750	749	111	<800	8682	8666
214	<800	7644	7609	4438	43442
836	887	10084	9932	514	<800
1670	1651	3641	3589	3077	3072
2650	2609	7555	7609	1371	1337
1890	1872			4185	4087
4615	4522			719	<800
1321	1273			4130	4087

FEATURES
source

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP11-79K17"
/clone_lib="RP11 human BAC library 11"
BASE COUNT      37777 a 24752 c 24789 g 41265 t
ORIGIN

```

BASE COUNT
ORIGIN

Query Match	36.0%;	Score 40;	DB 9;	Length 128583;
Best Local Similarity	100.0%;	Pred. NO. 4.3e-12;		
Matches	40;	Conservative	0;	Mismatches 0;
			Indels	0;
			Gaps	0;

[illegible]

RESULT 6

LOCUS	POSITION	CHROMOSOME	CLONE	SEQUENCE
AC107979	148290 bp	DNA	linear	PRI 01-JUL-2002
AC107979	Homo sapiens	chromosome 15,	clone	CMD-3049M7, complete sequence.
AC107979				

SOURCE

ORGANISM	REFERENCE
Homo sapiens	Birren, B., Nussbaum, C., and Lander, E.
Eukaryota, Metazoa, Chordata, Craniata, Vertebrata, Euteleostomi	1 (bases 1 to 148290)
Mammalia, Eutheria, Primates, Catarrhini, Homnidae, Homo.	2 (bases 1 to 148290)
REFERENCE	
AUTHORS	
TITLE	
JOURNAL	
REFERENCE	
AUTHORS	

AUTHORS

Anderson, S., Barrna, N., Bastien, V., Boguslavsky, L., Boukhalter Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepey, Y., Colanabelo, M., Collins, S., Collymore, A., Cook, A., Cooke, P., DeArallano, K., Dewar, K., Diaz, J., S. Dodge, S., Faro, S. Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Glade, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Kerasas, A., Kells, C., Laroque, K., Lamazares, R., Landers, T., Lehoczy, J., Levine, R., Liang, R., Maclean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrum, J., Menus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Ntobu, C.

TITTLE
JOURN

REFERENCE
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 148290)
AUTHORS
Birnbaum R., Lindon J., Nusbaum C., Randor F., and S.
S.

AUTHORS

Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Plunkhant, P., Pierroni, N., Pollara, V., Raymond, C., Retter, R., Ribbeck, M., Riley, R., Rise, C., Rogov, P., Roman, J., Roselli, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stenge-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Triggillo, J., Tse, J., Vasiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, M.-J., Young, G., Zaimon, J., Zembek, L., Zimmer, A., and Zody, M. (2019) Submitted.

Submitted (24-JAN-2019)

Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 148290)
Birren, B., Linton, J., Nusbaum, C., Rand, E., All, J.

Birren, J.

Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,
Bouhagbeter, B., Brown, A., Camarata, J., Campodiano, A., Chang, J.,
Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collimore, A.,
Cook, A., Cooke, P., DeVellano, K., Dewar, K., Diaz, J. S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Fitzhugh, W., Gage, D.,
Galagan, J., Gardyna, S., Glinde, S., Gold, S., Goyette, M., Graham, L.,
Grand, Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I.,
Johnson, R., Jones, C., Kamat, A., Karakas, A., Kells, C., Larocque, K.,
Lammaras, R., Landers, T., Lehoczy, J., Levine, R., Lindblad-Toh, K.,
Lin, G., Maclean, C., MacDonald, P., Major, J., Marquis, N.,
Mathews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrum, J.,
Menuss, L., Milnova, T., Mienna, V., Murphy, T., Naylor, J., Nguyen, C.,
Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P.,
O'Neill, D., Oliver, J., Peterson, K., Phukthong, P., Pierre, N.,
Pollard, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C.,
Rogov, P., Roman, J., Roselli, M., Roy, A., Santos, R., Schauer, S.,
Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange, Thomann, N.,
Stojanovic, N., Strauss, N., Sudirmanian, A., Talamas, J., Testaye, S.,
Thedore, J., Topham, K., Travers, M., Travis, N., Trigglio, J.,
Vassiliev, H., Viel, R., Vo, A., Wilson, M., Wu, X., Wyman, D.,
Young, G., Zainoun, J., zempek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (20-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
(bases 1 to 148290)

Birren, B

Barna, N., Bittan, V., Bloom, T., Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Chazaro, B., Cheepel, Y., Collymore, A., Cook, A., Cooke, P., Detrellano, K., Dewar, K., Diaz, J., S., Dodge, S., Fato, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gaidyna, S., Gird, S., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., leaders, T., Levine, R., Lindblad-roh, K., Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrum, J., Menus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunhkan, P., Pierre, N., Raymond, C., Retter, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, N., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange, S., Stojanack, R., Stojanovic, N., Talamas, J., Tefsey, S., Theodore, J., Toppan, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted 101-TH

Submitted (19.05.2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 26, 2002 this sequence version replaced g1:213131840.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

COMMENT

All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WtBR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: L24533
Center clone name: 3049_M7


```

TITLE
JOURNAL
REFERENCE
AUTHORS
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Submitted (01-DEC-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 165649)
Birtten, B., Nusbaum, C., Lander, E., All, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhaltier, B.,
Camata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.,
Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hume, W., Iliev, I., Johnson, R., Jones, C., Kamt, A.,
Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., Maclean, C., Macdonald, P., Major, D., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Mhova, T., Mlenga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Testaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 165649)
Birtten, B., Nusbaum, C., Lander, E., All, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhaltier, B.,
Camata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.,
Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hume, W., Iliev, I., Johnson, R., Jones, C., Kamt, A.,
Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., Maclean, C., Macdonald, P., Major, D., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Mhova, T., Mlenga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Testaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 1, 2002 this sequence version replaced qt:21592191.
All repeats were identified using RepeatMasker:
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L21917
Center clone name: 76_E_17
-----
FEATURES
source
Location/Qualifiers
1..165649
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-76E17"
/clone_1kb="RP11-11 Human Male BAC"
/complement(2..865)
/rpt_family="L1PA13"
repeat_region
complement(864..894)

```

```

/rpt_family="L1PA13"
complement(895..1248)
/rpt_family="TRHE1A"
complement(1249..1504)
/rpt_family="L1PA13"
1505..1626
/rpt_family="AluXs"
1627..1659
/rpt_family="(CAA)n"
1660..1831
/rpt_family="AluXs"
complement(1832..3132)
/rpt_family="L1PA13"
complement(3140..3699)
/rpt_family="L1M4C"
complement(4461..4755)
/rpt_family="AluXs"
5027..5397
/rpt_family="L2"
complement(5650..6330)
/rpt_family="L1ME1"
complement(6343..6460)
/rpt_family="L1ME1"
7214..7319
/rpt_family="L1MC3"
7354..7655
/rpt_family="L1MC3"
7659..7743
/rpt_family="L1PA10"
complement(7745..7843)
/rpt_family="AluSp/q"
7844..14156
/rpt_family="L1PA10"
14198..14328
/rpt_family="AluY"
14340..14503
/rpt_family="(TA)n"
14507..15032
/rpt_family="L1MC3"
15036..15220
/rpt_family="L1MC3"
complement(15218..16449)
/rpt_family="L1PA4"
16450..18154
/rpt_family="L1PA4"
18155..19210
/rpt_family="L1MC3"
19211..19266
/rpt_family="(TA)n"
19267..19372
/rpt_family="L1MC3"
19401..19476
/rpt_family="(TTATA)n"
19483..19546
/rpt_family="(CATATA)n"
19596..19625
/rpt_family="AT-rich"
19696..19750
/rpt_family="GA-rich"
19752..19920
/rpt_family="L1MD3"
19994..20102
/rpt_family="L2"
20049..20114
/notes="single clone coverage"
20485..20655
/rpt_family="MIR3"
complement(21285..21441)
/rpt_family="MIR"
21496..21717
/rpt_family="L2"
complement(21720..21796)
/rpt_family="MIR"

```

```
repeat_region complement(21943. .22131)
/rpt_family="MIR"
23082. .23195
repeat_region /rpt_family="L2"
23198. .23248
repeat_region /rpt_family="GA-rich"
23267. .23333
repeat_region /rpt_family="(CAT)n"
complement(24465. .24833)
/rpt_family="MT1A2"
26142. .26334
repeat_region /rpt_family="MIR"
complement(27659. .27811)
repeat_region /rpt_family="MIR"
complement(28811. .28860)
repeat_region /rpt_family="MIR"
complement(29026. .29269)
repeat_region /rpt_family="MIR"
complement(29673. .29706)
repeat_region /rpt_family="MSTB"
```

Very Match	36.0%;	Score 40;	DB 9;	Length 165649;
Best Local Similarity	100.0%;	Pred. No. 4.3e-12;		
Matches 40;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

```

0y      68 TAAACCTCTGGGTCTCTGTGTGTCCTAGTGGCTGCTC 107
         |||||
Db      8110 TAAACCTCTGGGTCTCTGTGTGTCCTAGTGGCTGCTC 8071

```

RESULT 8	AC104303	AC104303	178650 bp	DNA	linear	PR1 25-FEB-2007
LOCUS	AC104303					
DEFINITION	Homo sapiens chromosome 3 clone RP11-331P4, complete sequence.					
ACCESSION	AC104303	AC064830				
VERSION	AC104303.2	GT:18874945				
KEYWORDS	HTG.					
SOURCE	Homo sapiens.					
ORGANISM	Homo sapiens.					

REFERENCE	1 (bases 1 to 178650)
AUTHORS	Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z., Saenphimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE	Direct Submission
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 178650)
AUTHORS	Kaul, R.K., Olson, M.V., Raymond, C. and Haugen, E.D.
TITLE	Direct Submission
JOURNAL	Submitted (07-DEC-2001) Genome Center, University of Washington Box 352145, Seattle, WA 98195, USA
REFERENCE	3 (bases 1 to 178650)
AUTHORS	Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z., Saenphimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE	Direct Submission
JOURNAL	Submitted (25-FEB-2002) Genome Center, University of Washington Box 352145, Seattle, WA 98195, USA
COMMENT	On Feb 25, 2002 this sequence version replaced g1:17402782.

```

Center: University of Washington Genome Center
Center Code: UWGC
Web site: http://www.genome.washington.edu
Contact: uwgchgs@u.washington.edu
Drafting Center: UWGSC
-----
Project Information
Center project name: chr-3
Center clone name: RP11-391P4 (bc0402)
-----
Summary statistics
Sequencing vector: unknown: 55% of reads
Sequencing vector: plasmid: 45% of reads
Chemistry: dye-terminator ET: 88% of reads
Chemistry: dye-terminator Bis Dye: 11% of reads
Assembly program: Phrap: version 0.990319

```

Consensus quality: 178494 bases at least Q40
Consensus quality: 178631 bases at least Q30
Consensus quality: 178650 bases at least Q20
Insert size: 178648; sum-of-contigs
Quality coverage: 8.0x in Q20 bases; sum-of-contigs

Overlapping Sequences:
5': RP11-475023 (UMGC:bc0439) AC023346
3': RP11-79K12 AC076969

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp. Base-by-base quality values are not generally visible from the Genbank flat file format but are available as part of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both inset and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400–800 bps) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

B9111

ECORI

HindIII

SedDerMap	FngPrnt	SedDerMap	FngPrnt	SedDerMap	FngPrnt
7050	7417	8696	8715	7163	7098
2067	2138	6	<800	6582	6518
13472	13264	3077	3083	512	<800
7644	7859	514	<800	449	<800
111	<800	4438	4377	7988	7884
798	783	8682	8715	11779	11698
1098	1109	1896	1876	472	<800
5288	5395	6724	6948	783	787
4649	4507	10961	10773	926	926
461	<800	3055	3083	2431	2538
967	982	470	<800	1015	1029
1288	1268	1066	1037	758	<800
3045	2923	321	<800	1550	1541
743	783	2722	2683	361	<800

5628	5682	1204	1300	763	<800
2714	2699	6992	7281	2975	3002
4002	3900	7210	7830	3299	3468
227	<800	7821	8101	1907	1894
1794	1784	6855	6948	4020	4220
2168	2138	859	857	5158	5126
10032	9744	2629	2683	497	<800
6694	6610	1065	1037	4321	4220
1505	1491	3934	3965	887	926
3312	3304	859	857	180	<800
896	912	931	935	4969	4887
5873	5938	858	857	4755	4674
78	<800	799	857	26	<800
2135	2138	1895	1876	906	926
133	<800	58	<800	2325	2346
11939	11575	5129	5076	369	<800
1507	1491	6051	6048	2523	2538
8069	8181	3538	3537	1889	1894
1632	1617	1114	1094	4437	4428
3729	3678	8420	8391	373	<800
5553	5395	1024	1037	24	<800
358	<800	1333	1380	263	<800
3764	3678	1502	1481	632	<800
7022	6810	4538	4539	1258	1247
8534	8821	1414	1380	1114	1083
5206	5050	521	<800	1370	1320
9052	9744	8346	8391	1262	1247
1625	1617	39	<800	1546	1541
749	783	2716	2683	459	<800
10190	9744	2546	2683	135	<800
3857	3795	1064	1037	3142	3183
6144	6490	926	935	109	<800
514	<800	5154	5076	7858	7884
1364	1342	1975	1981	1891	1894
669	<800	628	<800	723	<800
		2398	2399	6975	7098
		8148	8391	1878	1894

Query Match 36.0%, Score 40; DB 9; Length 178650;
 Best Local Similarity 100.0%; Pred. No. 4; 3e-12;
 Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 69 AAAACCTCGGCTCTGTGTGCTGAGTGGCTGCT 108
 Db 165449 AAACTCTGGGTCTGTGTGCTGAGTGGCTGCT 165488

RESULT 9 192826 bp DNA linear PRI 28-FEB-2002
 AC090762 Homo sapiens chromosome 15, clone RP11-387E8, complete sequence.
 LOCUS AC090762/c
 DEFINITION AC090762.9 GI:18991378
 ACCESSION AC090762.9
 VERSION AC090762.9
 KEYWORDS HTG.
 SOURCE Homo sapiens.
 ORGANISM Homo sapiens.

REFERENCE 1 (bases 1 to 192826)
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 TITLE Homo sapiens chromosome 15, clone RP11-387E8
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 192826)
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,
 Barina, N., Bastien, V., Boguslavsky, L., Bouckgalter, B., Brown, A.,
 Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S.,
 Collamore, A., Cooke, P., Dearrellano, K., Dewar, K., Diaz, J. S.,
 Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,
 Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
 Jones, C., Karatas, A., Laroque, K., Lamazares, R., Landers, T.,
 Lehoczy, J., Levine, R., Liu, G., Maclean, C., Macdonald, P.,
 Marquis, N., Mathews, C., McCarthy, M., McEwan, P., McKernan, K.,
 McPheters, R., Meldrum, J., Meneus, L., Mihova, T., Mlenga, V.,
 Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H.,
 O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
 Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R.,
 Riback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Roselli, M.,
 Roy, A., Santos, R., Schauer, S., Schupback, R., Seaman, S., Severy, P.,
 Sougnuez, C., Spencer, B., Strange-Thomann, N., Stojanovic, N.,
 Strausz, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,
 Travers, M., Travis, N., Triggillo, J., Vassiliev, H., Viel, R., Vo, A.,
 Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J.,
 Zembek, L., Zimmer, A. and Zody, M.

TITLE Direct Submission
 JOURNAL Submitted (10-MAR-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 REFERENCE 3 (bases 1 to 192826)

AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., All, A., Allen, N.,
 Anderson, S., Barina, N., Bastien, V., Boguslavsky, L., Bouckgalter, B.,
 Brown, A., Camarata, J., Campopiano, A., Chang, D., Chazaro, B.,
 Choepel, Y., Colangelo, M., Collins, S., Collamore, A., Cook, A.,
 Cooke, P., Dearrellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S.,
 Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S.,
 Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
 Karatas, A., Karatas, A., Kells, C., Laroque, K., Lamazares, R.,

TITLE
JOURNAL
COMMENT

Landers, T., Lehoczy, J., Levine, R., Liu, G., MacLean, C.,
Mackdonald, P., Major, J., Marguis, N., Matthews, C., McCarthy, M.,
McEwan, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T.,
Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,
Notman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,
Peterson, K., Phukhang, P., Pierre, N., Pollara, V., Raymond, C.,
Retta, R., Riback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,
Roselli, M., Roy, A., Santos, R., Schauer, S., Schuppach, R., Seaman, S.,
Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
Topham, K., Travers, M., Travis, N., Triggillo, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,
Zainoun, J., Zemdek, L., Zimmer, A. and Zody, M.

Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Feb 28, 2002 this sequence version replaced g1:18377189.
All repeats were identified using RepeatMasker:
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIRB
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu

FEATURES
Source

Location/Qualifiers
1. 192826
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-387B8"
/clone_lib="RPCT-11 Human Male BAC"
repeat_region
complement(864..1222)
/rpt_family="L1PB3"
1223..1252
complement(1253..1492)
/rpt_family="(TAGC)n"
repeat_region
complement(1253..1492)
/rpt_family="L1PB3"
4287..4576
/rpt_family="AluJb"
4671..5034
/rpt_family="THE1B"
5130..5438
/rpt_family="AluSx"
5517..5662
/rpt_family="MIR"
5798..5827
/rpt_family="AT-rich"
5985..6114
/rpt_family="AluJb"
6227..6253
/rpt_family="(CA)n"
7329..7446
/rpt_family="MIR"
complement(8147..8452)
/rpt_family="AluJo"
8540..8639
/rpt_family="MER45"
complement(8832..8916)
/rpt_family="L2"
10391..10599
/rpt_family="MER3"
11688..11688
/rpt_family="L1MC/D"
12029..12078
/rpt_family="AT-rich"
12092..12447
/rpt_family="THE1C"

repeat_region complement(13616..13751)
/rpt_family="MIR3"
repeat_region complement(13958..14142)
/rpt_family="MIR"
repeat_region complement(15277..15553)
/rpt_family="MER8"
repeat_region complement(15811..16005)
/rpt_family="MIR"
repeat_region complement(16101..16440)
/rpt_family="L3"
16920..16958
/rpt_family="(TCCC)n"
complement(17145..17444)
/rpt_family="AluSx"
complement(18418..19953)
/rpt_family="L1MEC"
complement(19978..20262)
/rpt_family="L1MEC"
complement(20288..20794)
/rpt_family="L1MEC"
complement(20822..21097)
/rpt_family="L1MEC"
complement(21345..21743)
/rpt_family="L1MEC"
22599..23518
/rpt_family="L1MEC"
23527..23901
/rpt_family="L1MEC"
complement(23927..24026)
/rpt_family="MSTR1"
complement(24027..24256)
/rpt_family="MER30"
complement(24257..24557)
/rpt_family="MSTR1"
24563..24594
/rpt_family="AT-rich"
complement(25258..25639)
/rpt_family="L1MCc"
complement(25688..25838)
/rpt_family="L1MCc"
26167..26506
/rpt_family="THE1B"
26746..27094
/rpt_family="Tigger2a"
27095..27184
/rpt_family="MADE1"
27185..27279
/rpt_family="Tigger2a"
28321..28363
/rpt_family="(TG)n"
29172..29333
/rpt_family="MIR"
30664..30898
/rpt_family="L1MB8"
31577..31598
/rpt_family="AT-rich"
32378..32475
/rpt_family="CT-rich"
34655..34960
/rpt_family="AluY"
complement(35296..35389)
/rpt_family="MER5B"
complement(35419..35517)
/rpt_family="L1MC4"
complement(35518..35823)
/rpt_family="AluSx"
complement(35824..36030)
/rpt_family="L1MC4"
36682..36758
/rpt_family="MER5A"
36975..37160
/rpt_family="MER5A"
38140..38353

```

repeat_region      /rpt_family="MER96B"
                   38633..38656
repeat_region      /rpt_family="(TTTA)n"
                   complement(38657..40056)
repeat_region      /rpt_family="LIPAS"
                   complement(40060..40250)

Query Match
Best Local Similarity 100.0%; Score 40; DB 9; Length 192826;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 68 TAAACTCTGGCTCTGTGTGCTGAGTGCTGCTC 107
      |||
Db 190946 TAAACTCTGGCTCTGTGTGCTGAGTGCTGCTC 190907

RESULT 10
AC007445      32918 bp      DNA      linear      HTG 30-JUN-2000
US            Homo sapiens chromosome 18 clone RP11-344B7 map 18, *** SEQUENCING
AC007445      IN PROGRESS *** 1 ordered piece.
AC007445      HTG: HTGS_P1844149
KEYWORDS      Homo sapiens.
SOURCE        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
ORGANISM      Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE      1 (bases 1 to 32918)
AUTHORS      Birren, B., Linton, L., Nusbaum, C. and Lander, E.
JOURNAL      Homo sapiens chromosome 18, clone RP11-344B7
TITLE        Unpublished
COMMENT      2 (bases 1 to 32918)
              Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, M.,
              Baker, J., Baldwin, J., Barna, N., Beckey, R., Berr, J., Brown, A.,
              Castle, A., Cerny, J., Colangelo, M., Collins, S., Collymore, A.,
              Cooke, P., Dekrellano, K., Depayre, E., Devon, K., Dewar, K.,
              Donelan, L., Doyle, M., Ferrel, P., FitzHugh, W., Forrest, C.,
              Funke, R., Gage, D., Galagan, J., Gardy, S., Gilbert, D., Grant, G.,
              Hagos, B., Heaford, A., Horton, L., Howland, J. C., Jones, C., Kann, L.,
              Karats, A., Lehoczy, J., Lieu, C., Locke, K., Macdonald, P.,
              Marquis, N., McEwan, P., McKus, A., McKernan, K., McLaughlin, J.,
              Meidrum, J., Molla, M., Morris, W., Morrow, J., Mychaleckyj, J.,
              Naylor, J., Nilot, M., O'Connor, T., O'Donnell, P., Pavlin, B.,
              Peterson, K., Pollara, V., Riley, R., Roberts, D., Roy, A., Severy, P.,
              Stange-Thomann, N., Stojanovic, N., Stone, C., Subramanian, A.,
              Testaye, S., Torruella-Miller, I., Vassiliev, H., Vo, A., Wagner, A.,
              Wheeler, J., Wu, X., Wyman, D., Ye, W. J. and Zody, M.
              Direct Submission
              Submitted (30-APR-1999) Whitehead Institute/MIT Center for Genome
              Research, 320 Charles Street, Cambridge, MA 02141, USA
              On Jun 30, 2000 this sequence version replaced g1:8705092.
              All repeats were identified using RepeatMasker:
              Smit, A. F. A. & Green, P. (1996-1997)
              http://ftp.genome.washington.edu/RM/RepeatMasker.html

              Genome Center
              Center: Whitehead Institute/MIT Center for Genome Research
              Center code: WIBR
              Web site: http://www-seq.wi.mit.edu
              Contact: sequence_submissions@genome.wi.mit.edu
              Project Information
              Center project name: L571
              Center clone name: 344_B_7

* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

```

```

FEATURES
source
1 32918: contig of 32918 bp in length.
Location/Qualifiers
1..32918
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18"
/clone="RP11-344B7"
/clone_lib="RPC1-11 Human Male BAC"
BASE COUNT 9462 a 6493 c 6865 g 9853 t 245 others
ORIGIN
Query Match
Best Local Similarity 100.0%; Score 39; DB 2; Length 32918;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAACCTCTGCTGTGTGCTGAG 98
      |||
Db 3535 GGAGTATGTAACCTCTGCTGTGTGCTGAG 3573

RESULT 11
AL358817      38936 bp      DNA      linear      PRI 06-OCT-2001
LOCUS        Human DNA sequence from clone RP11-399N22 on chromosome 10,
DEFINITION   complete sequence.
ACCESSION    AL358817
VERSION      AL358817.18 GI:15990637
KEYWORDS     HTG.
SOURCE       human.
ORGANISM     Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE     1 (bases 1 to 38936)
AUTHORS      Lovell, J.
JOURNAL      Direct Submission
              Submitted (06-OCT-2001) Sanger Centre, Hinxton, Cambridgeshire,
              CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
              On Oct 9, 2001 this sequence version replaced g1:14669268.
              requests: clonerequests@sanger.ac.uk
              During sequence assembly data is compared from overlapping clones.
              Where differences are found these are annotated as variations
              together with a note of the overlapping clone name. Note that the
              variation annotation may not be found in the sequence submission
              corresponding to the overlapping clone, as we submit sequences with
              only a small overlap as described above.
              This sequence was finished as follows unless otherwise noted: all
              regions were either double-stranded or sequenced with an alternate
              chemistry or covered by high quality data (i.e., phred quality >=
              30); an attempt was made to resolve all sequencing problems, such
              as compressions and repeats; all regions were covered by at least
              one plasmid subclone or more than one M13 subclone; and the
              assembly was confirmed by restriction digest. The following
              abbreviations are used to associate primary accession numbers given
              in the feature table with their source databases: Em., EMBL; SW.,
              SWISSPROT; Tr., TrEMBL; Wp., WormPEP; Information on the WormPEP
              database can be found at
              http://www.sanger.ac.uk/Projects/C_elegans/wormpep
              This sequence
              was generated from part of bacterial clone contigs of human
              chromosome 10, constructed by the Sanger Centre Chromosome 10
              Mapping Group. Further information can be found at
              http://www.sanger.ac.uk/HGP/Chr10
              RP11-399N22 is from the library RPC1-11.2 constructed by the group
              of Pieter de Jong. For further details see
              http://www.chori.org/bacpac/home.htm
              VECTOR: pBAC3.6
              IMPORTANT: This sequence is not the entire insert of clone
              RP11-399N22. It may be shorter because we sequence overlapping
              sections only once, except for a short overlap.
              The true left end of clone RP11-43299 is at 36937 in this sequence.
              The true right end of clone RP11-91A1 is at 2000 in this sequence.
              Location/Qualifiers
              1..38936

```

FEATURES
source

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-399N22"
/clone_lib="RPCT-11.2"
/clone_lib="RPCT-11.2"
BASE COUNT 9315 a 9079 c 9111 g 11431 t
ORIGIN

Query Match 35.1%; Score 39; DB 9; Length 38936;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAACCTCTGGTCTCTGTGTGCTGAG 98
|||||
Db 28896 GGAGTATGTAACCTCTGGTCTCTGTGTGCTGAG 28934

RESULT 12
AC025179 124271 bp DNA linear HTG 20-APR-2001
LOCUS Homo sapiens chromosome 5 clone CTD-2174B5, WORKING DRAFT SEQUENCE,
DEFINITION 8 unordered pieces.
AC025179
VERSION AC025179.4 GI:13699647
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFN.
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS 1 (bases 1 to 124271)
TITLE DOE Joint Genome Institute.
JOURNAL Sequencing of Human Chromosome 5
AUTHORS 2 (bases 1 to 124271)
REFERENCE Unpublished
TITLE DOE Joint Genome Institute.
JOURNAL Direct Submission
AUTHORS Submitted (07-MAR-2000) Production Sequencing Facility, DOE Joint
TITLE Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
JOURNAL On Apr 20, 2001 this sequence version replaced gi:7711794.
COMMENT -----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov

Project Information
Center Project Name: 694394
Center clone name: CITB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1052: contig of 1052 bp in length
* 1053 1152: gap of unknown length
* 3331 3331: contig of 2179 bp in length
* 3332 3431: gap of unknown length
* 3432 7370: contig of 3939 bp in length
* 7371 7470: gap of unknown length
* 7471 15483: contig of 8013 bp in length
* 15484 15583: gap of unknown length
* 15584 24916: contig of 9333 bp in length

* 24917 25016: gap of unknown length
* 25017 39922: contig of 1490 bp in length
* 39923 40022: gap of unknown length
* 40023 68684: contig of 28662 bp in length
* 68685 68784: gap of unknown length
* 68785 124271: contig of 55487 bp in length.
Location/Qualifiers
1. 124271
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2174B5"
/clone_lib="Caltech human BAC library D"
BASE COUNT 36863 a 24233 c 23599 g 38876 t 700 others
ORIGIN

Query Match 35.1%; Score 39; DB 2; Length 124271;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 99
|||||
Db 79646 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 79684

RESULT 13
AC008814 146671 bp DNA linear PRI 31-OCT-2001
LOCUS Homo sapiens chromosome 5 clone CTD-2117L12, complete sequence.
DEFINITION AC008814
AC008814
VERSION AC008814.6 GI:16554342
KEYWORDS HTG.
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
TITLE 1 (bases 1 to 146671)
JOURNAL Direct Submission
AUTHORS DOE Joint Genome Institute and Stanford Human Genome Center.
TITLE Direct Submission
JOURNAL Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
AUTHORS Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
JOURNAL 3 (bases 1 to 146671)
REFERENCE DOE Joint Genome Institute and Stanford Human Genome Center.
TITLE Direct Submission
JOURNAL Submitted (31-OCT-2001) DOE Joint Genome Institute, 2800 Mitchell
AUTHORS Drive, Walnut Creek, CA 94598, USA
JOURNAL On Oct 31, 2001 this sequence version replaced gi:15290309.
COMMENT Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing Completed at Stanford Human Genome Center
www.shgc.stanford.edu
Quality: Phrap Quality >=40 99.3% of Sequence;
Estimated Total Number of Errors is 0.7.
Location/Qualifiers
1. 146671
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2117L12"
BASE COUNT 44951 a 28815 c 28434 g 44471 t
ORIGIN

Query Match 35.1%; Score 39; DB 9; Length 146671;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 99
|||||
Db 54480 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 54518

RESULT 14
AP001019/c
LOCUS Homo sapiens chromosome 18 clone RP11-752111 map 18p11.3, WORKING
DEFINITION DRAFT SEQUENCE, 24 unordered pieces.
ACCESSION AP001019
VERSION AP001019.2 GI:8117689
KEYWORDS HTG: HTGS_PHASE1: HTGS_DRAFT.
SOURCE Homo sapiens DNA, clone:RP11-752111.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 159747)
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Matanabe, H. and Sakaki, Y.
TITLE Homo sapiens 159,747 genomic DNA of 18p11.3
2 (bases 1 to 159747)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Matanabe, H. and Sakaki, Y.
DIRECT SUBMISSION
Submitted (05-JAN-2000) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
Kilasato Univ., 1-15-1 Kilasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gs.c.riken.go.jp,
URL:http://hgp.gs.c.riken.go.jp/, Tel:01-42-778-9923,
Fax:01-42-778-9924)
On May 31, 2000 this sequence version replaced gi:6997769.
COMMENT
----- Genome Center
Center: RIKEN Genomic Sciences Center(GSC)
Center code: RIKEN
Web site: http://hgp.gs.c.riken.go.jp/
Contact: hattori@gs.c.riken.go.jp
----- Project Information
Center project name: Humdraft18
Center clone name: RP11-752111
----- Summary Statistics
Sequencing vector: PCR products; 100% of reads
Chemistry: Dye-terminator ET-amersham; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 145356 bases at least Q40
Consensus quality: 152227 bases at least Q30
Consensus quality: 155843 bases at least Q20
Insert size: 157447; sum-of-contigs
Quality coverage: 4.51x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of
24 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs 'N', but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved
1 18444 contig of 18444 bp in length
18545 29631 contig of 11087 bp in length
29732 43143 contig of 13412 bp in length
43244 57170 contig of 13927 bp in length
57271 68240 contig of 10970 bp in length
68341 78134 contig of 9794 bp in length
78235 87873 contig of 9639 bp in length
87974 96197 contig of 8224 bp in length
96298 103976 contig of 7679 bp in length
104077 110726 contig of 6650 bp in length
110827 116866 contig of 5823 bp in length
116967 122789 contig of 5823 bp in length
122890 127665 contig of 4776 bp in length
127766 132618 contig of 4853 bp in length
132619 13718: gap of 100 bp
13719 136051 contig of 3333 bp in length
136052 136151: gap of 100 bp
136152 139749: contig of 3598 bp in length
139750 139849: gap of 100 bp
139850 142810: contig of 2861 bp in length
142811 142910: gap of 100 bp
142911 145855: contig of 2945 bp in length
145856 145955: gap of 100 bp
145956 148840: contig of 2885 bp in length
148841 148940: gap of 100 bp
148941 151243: contig of 2203 bp in length
151244 151343: gap of 100 bp
151344 153454: contig of 2111 bp in length
153455 153554: gap of 100 bp
153555 156134: contig of 2580 bp in length
156135 156234: gap of 100 bp
156235 158089: contig of 1855 bp in length
158090 158189: gap of 100 bp
158190 159747: contig of 1558 bp in length.
Location/Qualifiers
1. 159747
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18p11.3"
/clone="RP11-752111"
1. 18444
/note="assembly_fragment"
18545.29631
/note="assembly_fragment"

145956 148840 contig of 2885 bp in length
148941 151243 contig of 2303 bp in length
151344 153454 contig of 2111 bp in length
153555 156134 contig of 2580 bp in length
156235 158089 contig of 1855 bp in length
158190 159747 contig of 1558 bp in length
Sequence updated (26-May-2000)
* NOTE: This is a 'working draft' sequence. It currently
* consists of 24 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of 'N', but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 18444: contig of 18444 bp in length
18445 18544: gap of 100 bp
18545 29631: contig of 11087 bp in length
29632 29731: gap of 100 bp
29732 43143: contig of 13412 bp in length
43144 43243: gap of 100 bp
43244 57170: contig of 13927 bp in length
57171 57270: gap of 100 bp
57271 68240: contig of 10970 bp in length
68241 68340: gap of 100 bp
68341 78134: contig of 9794 bp in length
78135 78234: gap of 100 bp
78235 87873: contig of 9639 bp in length
87874 87973: gap of 100 bp
87974 96197: contig of 8224 bp in length
96198 96297: gap of 100 bp
96298 103976: contig of 7679 bp in length
103977 104076: gap of 100 bp
104077 110726: contig of 6650 bp in length
110727 110826: gap of 100 bp
110827 116866: contig of 6040 bp in length
116867 116966: gap of 100 bp
116967 122789: contig of 5823 bp in length
122790 122889: gap of 100 bp
122890 127665: contig of 4776 bp in length
127666 127765: gap of 100 bp
127766 132618: contig of 4853 bp in length
132619 13718: gap of 100 bp
13719 136051: contig of 3333 bp in length
136052 136151: gap of 100 bp
136152 139749: contig of 3598 bp in length
139750 139849: gap of 100 bp
139850 142810: contig of 2861 bp in length
142811 142910: gap of 100 bp
142911 145855: contig of 2945 bp in length
145856 145955: gap of 100 bp
145956 148840: contig of 2885 bp in length
148841 148940: gap of 100 bp
148941 151243: contig of 2203 bp in length
151244 151343: gap of 100 bp
151344 153454: contig of 2111 bp in length
153455 153554: gap of 100 bp
153555 156134: contig of 2580 bp in length
156135 156234: gap of 100 bp
156235 158089: contig of 1855 bp in length
158090 158189: gap of 100 bp
158190 159747: contig of 1558 bp in length.
Location/Qualifiers
1. 159747
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18p11.3"
/clone="RP11-752111"
1. 18444
/note="assembly_fragment"
18545.29631
/note="assembly_fragment"

```
misc_feature 29732..43143
              /note="assembly-fragment"
misc_feature 43244..57170
              /note="assembly-fragment clone_end:SP6 vector_side:left"
misc_feature 57271..68240
              /note="assembly-fragment"
misc_feature 68341..78134
              /note="assembly-fragment"
misc_feature 78235..87873
              /note="assembly-fragment"
misc_feature 87974..96197
              /note="assembly-fragment"
misc_feature 96298..103976
              /note="assembly-fragment clone_end:77 vector_side:left"
misc_feature 104077..110726
              /note="assembly-fragment"
misc_feature 110827..116866
              /note="assembly-fragment"
misc_feature 116867..122789
              /note="assembly-fragment"
misc_feature 122890..127665
              /note="assembly-fragment"
misc_feature 127766..132618
              /note="assembly-fragment"
misc_feature 132719..136051
              /note="assembly-fragment"
misc_feature 136152..139749
              /note="assembly-fragment"
misc_feature 139850..142810
              /note="assembly-fragment"
misc_feature 142811..145855
              /note="assembly-fragment"
misc_feature 145956..148840
              /note="assembly-fragment"
misc_feature 148941..151243
              /note="assembly-fragment"
misc_feature 151344..153454
              /note="assembly-fragment"
misc_feature 153555..156134
              /note="assembly-fragment"
misc_feature 156235..158089
              /note="assembly-fragment"
misc_feature 158190..159747
              /note="assembly-fragment"
BASE COUNT 47302 a 30397 c 29897 g 49849 t 2302 others
ORIGIN
Query Match 35.1%: Score 39; DB 2; Length 159747;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
60 GGAGTATGTAACCTCCTGGCTCTGTGTGCTGAG 98
|||||
DB 74932 GGAGTATGTAACCTCCTGGCTCTGTGTGCTGAG 74894
|||||
RESULT 15
AC034249 162740 bp DNA linear HTG AC034249
LOCUS Homo sapiens chromosome 5 clone RP11-427C17, WORKING DRAFT
DEFINITION
SEQUENCE, 6 ordered pieces.
ACCESSION AC034249
VERSION AC034249.3 GI:15383785
KEYWORDS HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 162740)
AUTHORS DOE Joint Genome Institute.
TITLE Sequencing of Human Chromosome 5
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 162740)
```

```
AUTHORS DOE Joint Genome Institute.
TITLE Direct Submission
JOURNAL Submitted (05-APR-2000) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
COMMENT On Aug 31, 2001 this sequence version replaced gi:9211234.
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 570398
Center clone name: RPCI-11_427C17
-----
Summary Statistics
Consensus quality: 158733 bases at least Q40
Consensus quality: 161469 bases at least Q30
Consensus quality: 162087 bases at least Q20
Estimated insert size: 160000; pulse field gel estimation
Estimated insert size: 162240; sum-of-contigs estimation
Quality coverage: 9.16 in Q20 bases; pulse field gel estimation
Quality coverage: 9.16 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
1 85871: contig of 85871 bp in length
85872 96765: contig of 10794 bp in length
96766 96865: gap of unknown length
96866 103302: contig of 6437 bp in length
103303 103402: gap of unknown length
103403 111971: contig of 8569 bp in length
111972 112071: gap of unknown length
112072 129034: contig of 16963 bp in length
129035 129134: gap of unknown length
129135 162740: contig of 33606 bp in length.
*
FEATURES
Location/Qualifiers
source
1..162740
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="RP11-427C17"
/clone_lib="RPCI human BAC library 11"
BASE COUNT 48722 a 31435 c 31563 g 50520 t 500 others
ORIGIN
Query Match 35.1%: Score 39; DB 2; Length 162740;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
61 GAGTATGTAACCTCCTGGCTCTGTGTGCTGAGT 99
|||||
DB 3599 GAGTATGTAACCTCCTGGCTCTGTGTGCTGAGT 3637
|||||
Search completed: April 25, 2003, 00:30:53
Job time : 719.885 secs
```


GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:19:38 ; Search time 51.0118 Seconds

(without alignments)
4900.271 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgagctcttgcctt.....gcctgagtcgctcctact 111

Scoring table: OLIGO-NUC
Gapop 60.0 , Gapext 60.0

Searched: 2185239 segs, 1125999159 residues

Read size: 0

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

N_Geneseq_101002:*

- 1: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1980.DAT:*
- 2: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1981.DAT:*
- 3: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1982.DAT:*
- 4: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1983.DAT:*
- 5: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1984.DAT:*
- 6: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1985.DAT:*
- 7: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1986.DAT:*
- 8: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1987.DAT:*
- 9: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1988.DAT:*
- 10: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1989.DAT:*
- 11: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1990.DAT:*
- 12: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1991.DAT:*
- 13: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1992.DAT:*
- 14: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1993.DAT:*
- 15: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1994.DAT:*
- 16: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1995.DAT:*
- 17: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1996.DAT:*
- 18: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1997.DAT:*
- 19: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1998.DAT:*
- 20: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1999.DAT:*
- 21: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2000.DAT:*
- 22: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
- 23: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
- 24: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	447	21 AAC03794	Human secreted pro
2	111	100.0	447	21 AAZ42680	Human 5' EST isola
3	24	21.6	660	23 AAS73441	DNA encoding novel
4	20	18.0	2526	23 AAS66674	DNA encoding novel
5	18	16.2	431	24 ABR24383	DNA encoding human
6	18	16.2	431	24 ABR24390	DNA encoding human
7	18	16.2	570	22 ABR63453	Human foetal liver
8	18	16.2	570	22 ABR30652	Probe #9118 for ge
9	18	16.2	570	22 AAK11985	Human brain expres

10	18	16.2	570	22 AAK37688	Human bone marrow
11	18	16.2	570	22 AAI18447	Probe #8380 for ge
12	18	16.2	570	22 AAI43563	Probe #12249 used
13	18	16.2	570	24 ABA11680	Human genome-deliv
14	18	16.2	827	23 AAS87115	DNA encoding novel
15	18	16.2	3342	23 AAS87118	DNA encoding novel
16	17	15.3	266	20 AAV89709	EST clone CT857.
17	17	15.3	531	23 ABR43012	Genomic sequence #
18	17	15.3	609	24 ABR80793	Bacillus clausii q
19	17	15.3	828	22 AAI94407	Human neuroblastom
20	17	15.3	1047	24 AAS62681	CDNA sequence #468
21	17	15.3	1363	9 AAN70128	Novel DNA encoding
22	17	15.3	1546	22 AAK82125	Human immune/haema
23	17	15.3	1982	21 AAC68089	Human secreted pro
24	17	15.3	6928	22 ABA21109	Human nervous syst
25	17	15.3	21340	23 ABL12924	Drosophila melanog
26	17	15.3	62909	22 AAF28545	Genomic fragment #
27	17	15.3	172637	24 ABR83124	Human voltage-actl
28	17	15.3	495289	24 ABO67195	Listeria innocua c
29	17	15.3	1503900	22 AAK95240	Human neuroguilin-1
30	17	15.3	1503900	22 AAK96733	Human neuroguilin-1
31	17	15.3	3011208	24 ABO69245	Listeria innocua D
32	16	14.4	79	22 AAC89232	Human brain T. calc
33	16	14.4	250	16 AAT22213	Human gene signatu
34	16	14.4	260	24 ABR16202	Human OREX polynuc
35	16	14.4	281	24 ABL67351	Thyroid cancer rel
36	16	14.4	302	24 ABR78873	Human ORF3820 CDNA
37	16	14.4	347	24 ABR60208	Human cancer relat
38	16	14.4	348	22 AAF65225	Novel human polynu
39	16	14.4	360	22 AAF57925	Human immune/haema
40	16	14.4	377	24 ABL38444	Human colon tumour
41	16	14.4	407	23 ABR36584	Human prostate exp
42	16	14.4	414	22 AAI82407	Human polynucleoti
43	16	14.4	415	23 ABR07896	Human prostate exp
44	16	14.4	432	23 ABR36813	Human prostate exp
45	16	14.4	434	22 AAI79941	Human polynucleoti

ALIGNMENTS

RESULT 1	AAAC03794	standard; cDNA; 447 bp.
ID	AAAC03794	
XX	AAAC03794;	
AC	06-OCT-2000 (first entry)	
XX		
DT		
XX		
DE	Human secreted protein 5' EST, SEQ ID NO: 3792.	
XX		
KW	Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;	
KW	gene therapy; chromosome mapping; ss.	
XX		
OS	Homo sapiens.	
XX		
PN	EP1033401-A2.	
XX		
PD	06-SEP-2000.	
XX		
PF	21-FEB-2000; 2000EP-0200610.	
XX		
PR	26-FEB-1999; 99US-0122487.	
XX		
PA	(GEST) GENSET.	
XX		
PI	Dumas Milne Edwards J, Duclert A, Giordano J;	
XX		
WP	WPI: 2000-500381/45.	
DR	P-PSDB; AAG03788.	
XX		
PT	New nucleic acid that is a 5' expressed sequence tag (5' EST) for	
PT	obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for	

diagnostic, forensic, gene therapy and chromosome mapping procedures -
Claim 1: SEQ ID 3792; 71bp + CD-ROM; English.

The present sequence is one of a large number of 5' ESTs derived from cDNAs encoding secreted proteins. An ORF has been identified within the sequence. The 5' ESTs were prepared from total human RNAs or poly(A) RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of cDNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design expression and secretion vectors.

Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;

Query Match 100.0%; Score 111; DB 21; Length 447;
Best Local Similarity 100.0%; Pred. No. 3e-46; Mismatches 0; Gaps 0;
Matches 111; Conservative 0; Indels 0;

QY 1 ATGGGTGATCTTTGCTTGCAGATTCCTTTTCATCTTTGACGAGACTTCGGGCGC 60
DB 51 ATGGGTGATCTTTGCTTGCAGATTCCTTTTCATCTTTGACGAGACTTCGGGCGC 110
QY 61 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTCTACT 111
DB 111 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTCTACT 161

RESULT 2

AA242680
ID AA242680 standard; cDNA: 447 BP.

AC AA242680;

DT 01-FEB-2000 (first entry)

DE Human 5' EST isolated from a cDNA library SEQ ID NO:439.

Human: 5' EST; expressed sequence tag; secreted protein; diagnostics;
gene therapy; chromosome mapping; upstream regulatory sequence;
forensic; location; development; protein synthesis; stability;
regulation; identification; ss.

OS Homo sapiens.

PN WO953051-A2.

PD 21-OCT-1999.

PF 09-APR-1999; 99WO-IB00712.

PR 09-APR-1998; 98US-0057719.

PR 28-APR-1998; 98US-0069047.

PA (GEST) GENSET.

PI Dumas Milne Edwards J, Duclert A, Giordano J;

DR WPI; 2000-038446/03.

DR P-PSDB; AAY65066.

PT Novel secreted protein 5' expressed sequence tag sequences used in
diagnostic, forensic, gene therapy, and chromosome mapping procedures

PS Claim 1: Page 402; 837bp; English.

XX AA24265 to AA243075 represent novel 5' expressed sequence tag (EST)

sequences, corresponding to human secreted proteins. AAY64651 to
AA16538 represent the EST-related proteins corresponding to AA24265 to
AA43052. The 5' ESTs can be used for producing secreted human gene
products. They can be used to identify and isolate 5' untranslated
regions (UTRs) and upstream regulatory regions which control the
location, development stage, rate, and quantity of protein synthesis, as
well as stability of mRNA. The ESTs are also useful as probes for
chromosome mapping, and to obtain full length cDNA clones. The ESTs can
also be used in forensic procedures to identify individuals, or in
diagnostic procedures to identify individuals having genetic diseases
resulting from abnormal gene expression. The products may also be used in
gene therapy protocols. The nucleic acids encoding signal peptides can be
used for directing extracellular secretion of a polypeptide or the
insertion of a polypeptide into a membrane, or importing a polypeptide
into a cell. The proteins encoded by the EST sequences may be useful in
treating a variety of human conditions. Secreted proteins have
therapeutic value, and the identification of new secreted proteins is
valuable. AA24249 to AA24264 and AAY6464 to AAY64650 represent
sequences used in the exemplification of the present invention.

Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;

Query Match 100.0%; Score 111; DB 21; Length 447;
Best Local Similarity 100.0%; Pred. No. 3e-46; Mismatches 0; Gaps 0;
Matches 111; Conservative 0; Indels 0;

QY 1 ATGGGTGATCTTTGCTTGCAGATTCCTTTTCATCTTTGACGAGACTTCGGGCGC 60
DB 51 ATGGGTGATCTTTGCTTGCAGATTCCTTTTCATCTTTGACGAGACTTCGGGCGC 110
QY 61 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTCTACT 111
DB 111 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTCTACT 161

RESULT 3

AA573441/C
ID AA573441 standard; cDNA: 660 BP.

AC AA573441;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #9245.

Human: chromosome mapping; gene mapping; gene therapy; forensic;
food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEO INC.

PI Dzmanac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

DR P-PSDB; ABG09254.

PT New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity

PS Claim 1; SEQ ID NO 9245; 103bp; English.

XX

CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantifying a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.

Sequence 660 BP; 149 A; 127 C; 148 G; 236 T; 0 other;

Query Match 21.6%; Score 24; DB 23; Length 660;

Best Local Similarity 100.0%; Pred. No. 0.017;

Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 69 AAACCTCGGCTCTGTGTG 92
|||||

Db 656 AAACCTCGGCTCTGTGTG 633

RESULT 4
AAS6674

ID AAS6674 standard; cDNA: 2526 BP.

XX AAS6674;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #2478.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

OS WO200175067-A2.

XX 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

XX 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

DR P-PSDB; ABG02487.

XX New isolated polynucleotide and encoded polypeptides, useful in

PT diagnostics, forensics, gene mapping, identification of mutations

PS Claim 1; SEQ ID No 2478; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and

CC polypeptide (II) sequences. (I) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantifying a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.

Sequence 2526 BP; 871 A; 536 C; 541 G; 578 T; 0 other;

Query Match 18.0%; Score 20; DB 23; Length 2526;

Best Local Similarity 100.0%; Pred. No. 1.7;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 TTTCATCTTTCGAGGACTT 51
|||||

Db 2377 TTTCATCTTTCGAGGACTT 2396

RESULT 5
ABK24383

ID ABK24383 standard; cDNA: 431 BP.

XX ABK24383;

DT 09-APR-2002 (first entry)

DE DNA encoding human lung cancer protein, Seq ID No 80.

XX Human; lung cancer; cytostatic; vaccine; gene; ss.

XX Homo sapiens.

PN WO200192525-A2.

PD 06-DEC-2001.

XX 25-MAY-2001; 2001WO-US17066.

PF 26-MAY-2000; 2000US-207485P.

PR 06-SEP-2000; 2000US-230475P.

XX (CORI-) CORIXA CORP.

PI Harlocker SL, Wang T, Bangur CS, Klee JT, Switzer A;

XX WPI; 2002-122068/16.

XX New lung tumour polypeptides and polynucleotides, useful in

PT pharmaceutical compositions, such as vaccines, for treating or

PS Claim 9; Page 172; 179pp; English.

XX The invention relates to novel human lung cancer polynucleotide (I)

CC and polypeptides (II). (I) and (II) are useful in pharmaceutical

CC compositions, such as vaccines, for the diagnosis and treatment of lung

CC cancer. The polynucleotides are also useful as probes or primers for

CC nucleic acid hybridisation. ABK24314-ABK24397 represent human lung

CC cancer coding sequences of the invention.

XX SQ Sequence 431 BP; 108 A; 97 C; 123 G; 101 T; 2 other;
Query Match 16.2%; Score 18; DB 24; Length 431;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
DB 18 TGCCTGAGTGCTGCTCT 35
RESULT 6
ABK24390
ID ABK24390 standard; CDNA; 431 BP.
XX
AC ABK24390;
XX
DT 09-APR-2002 (first entry)
XX
XX DNA encoding human lung cancer protein, Seq ID No 87.
XX
XX Human; lung cancer; cytostatic; vaccine; gene; ss.
XX
XX Homo sapiens.
XX
XX WO200192525-A2.
XX
XX PD 06-DEC-2001.
XX
XX PF 25-MAY-2001; 2001WO-US17066.
XX
XX PR 26-MAY-2000; 2000US-207485P.
XX
XX PR 06-SEP-2000; 2000US-230475P.
XX
XX PA (CORI-) CORIXA CORP.
XX
XX PI Harlocker SL, Wang T, Bangur CS, Klee JT, Switzer A;
XX
XX WPI: 2002-122068/16.
XX
XX DR New lung tumour polypeptides and polynucleotides, useful in
XX
XX PT pharmaceutical compositions, such as vaccines, for treating or
XX
XX PT preventing lung cancer, or as probes or primers for nucleic acid
XX
XX PT hybridisation -
XX
XX PS Claim 1; Page 174; 179pp; English.
XX
XX The invention relates to novel human lung cancer polynucleotide (I)
XX and polypeptides (II). (I) and (II) are useful in pharmaceutical
XX compositions, such as vaccines, for the diagnosis and treatment of lung
XX cancer. The polynucleotides are also useful as probes or primers for
XX nucleic acid hybridisation. ABK24314-ABK24397 represent human lung
XX cancer coding sequences of the invention.
XX
XX SQ Sequence 431 BP; 108 A; 97 C; 123 G; 101 T; 2 other;
Query Match 16.2%; Score 18; DB 24; Length 431;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
DB 18 TGCCTGAGTGCTGCTCT 35
RESULT 7
ABA63453/c
ID ABA63453 standard; DNA; 570 BP.
XX
AC ABA63453;
XX
XX
XX 01-FEB-2002 (first entry)

XX DE Human foetal liver single exon nucleic acid probe #11758.
XX
XX KW Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
XX
XX OS Homo sapiens.
XX
XX PN WO200157277-A2.
XX
XX PD 09-AUG-2001.
XX
XX PF 30-JAN-2001; 2001WO-US00669.
XX
XX PR 04-FEB-2000; 2000US-0180312.
XX
XX PR 26-MAY-2000; 2000US-0207456.
XX
XX PR 30-JUN-2000; 2000US-0608408.
XX
XX PR 03-AUG-2000; 2000US-0632366.
XX
XX PR 21-SEP-2000; 2000US-0234687.
XX
XX PR 27-SEP-2000; 2000US-0236359.
XX
XX PR 04-OCT-2000; 2000GB-0024263.
XX
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX DR WPI: 2001-483447/52.
XX
XX PT Human genome-derived single exon nucleic acid probes useful for
XX
XX PT analyzing gene expression in human fetal liver -
XX
XX PS Claim 1; SEQ ID NO 11758; 639pp + sequence listing; English.
XX
XX CC The invention relates to a single exon nucleic acid probe for
XX
XX CC measuring human gene expression in a sample derived from human foetal
XX
XX CC liver. The single exon nucleic acid probes may be used for predicting,
XX
XX CC measuring and displaying gene expression in samples derived from human
XX
XX CC fetal liver. The present sequence is a single exon nucleic acid
XX
XX CC probe of the invention.
XX
XX CC Note: The sequence data for this patent did not form part of the
XX
XX CC printed specification, but was obtained in electronic format directly
XX
XX CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
XX SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
Query Match 16.2%; Score 18; DB 22; Length 570;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 85 TGTGTGCTGCTGAGTGGC 102
|||||
DB 505 TGTGTGCTGCTGAGTGGC 488
RESULT 8
ABA30652/c
ID ABA30652 standard; DNA; 570 BP.
XX
AC ABA30652;
XX
XX
XX DT 23-JAN-2002 (first entry)
XX
XX DE Probe #9118 for gene expression analysis in human heart cell sample.
XX
XX DE Human; gene expression; heart; microarray; vascular system; probe;
XX
XX KW cardiovascular disease; hypertension; cardiac arrhythmia.
XX
XX KW congenital heart disease; ss.
XX
XX OS Homo sapiens.
XX
XX XX
XX PN WO200157274-A2.
XX
XX PD 09-AUG-2001.

PF 30-JAN-2001; 2001WO-US00666.
XX
PR 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488990/53.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
XX hearts -
PS Claim 1; SEQ ID No 9118; 530pp; English.

CC The present invention relates to single exon nucleic acid probes for
CC measuring human gene expression in a sample derived from human heart. The
CC present sequence is one such probe. The probes may be used for
CC predicting, measuring and displaying gene expression in samples derived
CC from the human heart via microarrays. By measuring gene expression, the
CC probes are useful for predicting, diagnosing, grading, staging,
CC monitoring and prognosing diseases of the human heart and vascular system
CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
CC congenital heart disease.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 85 TGTGTGTGCTGAGTGGC 102
XXXXXXXXXXXXXXXXXXXX
Db 505 TGTGTGTGCTGAGTGGC 488

RESULT 9
AAK1985/c
ID AAK1985 standard; DNA; 570 BP.
AAK1985;

DT 05-NOV-2001 (first entry)
XX
XX Human brain expressed single exon probe SEQ ID NO: 11976.
DE
XX
KW Human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
XX Homo sapiens.
OS
XX
PN WO200157275-A2.
PD
XX 09-AUG-2001.
PD
XX
PF 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-483446/52.
XX
XX Single exon nucleic acid probes for analyzing gene expression in human
XX brains -
PT
XX
PS Example 4; SEQ ID NO: 11976; 650pp + Sequence Listing; English.
XX
XX The present invention provides a number of single exon nucleic acid
XX probes which are derived from genomic sequences expressed in the human
XX brain. They can be used to measure gene expression in brain cell samples,
XX which may enable the diagnosis and improved treatment of nervous system
XX diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
XX epilepsy and cancers. The present sequence is one of the probes of the
XX invention.
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 85 TGTGTGTGCTGAGTGGC 102
XXXXXXXXXXXXXXXXXXXX
Db 505 TGTGTGTGCTGAGTGGC 488

RESULT 10
AAK37688/c
ID AAK37688 standard; DNA; 570 BP.
AAK37688;
XX
XX 06-NOV-2001 (first entry)
XX
XX Human bone marrow expressed single exon probe SEQ ID NO: 12245.
DE
XX
KW Human; bone marrow expressed exon; gene expression analysis; probe;
KW microarray; cancer; leukemia; lymphoma; myeloma; ss.
XX
XX Homo sapiens.
OS
XX
PN WO200157276-A2.
PD
XX 09-AUG-2001.
PD
XX
PF 30-JAN-2001; 2001WO-US00668.
XX
PR 04-FEB-2000; 2000US-0180312.
XX 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
XX 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
XX 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488990/53.
XX
XX Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human bone marrow -
PT
XX
PS Example 4; SEQ ID NO: 12245; 658pp + Sequence Listing; English.
XX
XX The present invention provides a number of single exon nucleic acid

CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers
CC such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC the probes of the invention.

XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGCC 102

Db 505 TGTGTGTCCTGAGTGCC 488

RESULT 11

AA118447/c

AA118447 standard; DNA: 570 BP.

AC AA118447;

DT 12-OCT-2001 (first entry)

DE Probe #8380 for gene expression analysis in human cervical cell sample.

XX Probe: human; microarray; gene expression; cervical epithelial cell;

KW cervical cancer; ss.

XX Homo sapiens.

OS WO200157278-A2.

PN 09-AUG-2001.

PD 30-JAN-2001; 2001WO-US00670.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PA Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-488901/53.

XX Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human cervical epithelial cells -

XX Claim 25; SEQ ID No 8380; 487pp; English.

CC The present invention relates to human single exon nucleic acid probes
CC (SENP). The present sequence is one such probe. The SENPs are derived
CC from human HeLa cells. The SENPs can be used to produce a single exon
CC microarray, which can be used for measuring human gene expression in a
CC sample derived from human cervical epithelial cells. By measuring gene
CC expression, the probes are therefore useful in grading and/or staging
CC of diseases of the cervix, notably cervical cancer.

CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGCC 102

Db 505 TGTGTGTCCTGAGTGCC 488

RESULT 12

AA143563/c

AA143563 standard; DNA: 570 BP.

AC AA143563;

DT 17-OCT-2001 (first entry)

DE Probe #12249 used to measure gene expression in human placenta sample.

XX Probe: microarray; human; placenta; antenatal diagnosis;

KW genetic disorder; ss.

XX Homo sapiens.

OS WO200157272-A2.

PN 30-JAN-2001; 2001WO-US00663.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PA Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-48897/53.

XX Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human placenta -

XX Claim 25; SEQ ID No 12249; 654pp; English.

CC The present invention relates to single exon nucleic acid probes (SENP).
CC The present sequence is one such probe. The probes are useful for
CC producing a microarray for predicting, measuring and displaying gene
CC expression in samples derived from human placenta. The probes are useful
CC for antenatal diagnosis of human genetic disorders.

XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGCC 102

Db 505 TGTGTGTCCTGAGTGCC 488

RESULT 13

ABSI1680/c

ABSI1680 standard; DNA: 570 BP.

DT 19-AUG-2002 (first entry)

DE Human genome-derived single exon probe from lung SEQ ID No 11671.

KW Human; ds; single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 XX
 OS Homo sapiens.
 PN WO200186003-A2.
 XX
 PD 15-NOV-2001.
 XX
 PF 30-JAN-2001; 2001WO-US000665.
 XX
 PR 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 DR WPI; 2002-114183/15.
 XX
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 PS Claim 1; SEQ ID No 11671; 634pp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridise at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray, assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a single exon
 CC probe of the invention.

CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 S0 Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
 Query Match 16.2%; Score 18; DB 24; Length 570;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 85 TGTGTGTCCTGAGTGC 102
 ||||||||||||||||
 Db 505 TGTGTGTCCTGAGTGC 488
 RESULT 14
 AAS87115
 ID AAS87115 standard; cDNA; 827 BP.
 XX
 AC AAS87115;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #22919.
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US008631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HSE-) HSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 DR P-PSDB; ABG22928.
 PT
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 1; SEQ ID No 22919; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed

CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 827 BP; 198 A; 202 C; 249 G; 178 T; 0 other;
Query Match 16.2%; Score 18; DB 23; Length 827;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
DB 344 TGCCTGAGTGCTGCTCT 361
RESULT 15
AAS87118/C
ID AAS87118 standard; cDNA; 3342 BP.
XX
PC AAS87118;
XX
13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #22922.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR MPI: 2001-639362/73.
DR P-PSDB; ABG22931.
XX
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID No 22922; 103pp; English.
XX
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX
SQ Sequence 3342 BP; 898 A; 796 C; 867 G; 781 T; 0 other;
Query Match 16.2%; Score 18; DB 23; Length 3342;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
DB 113 TGCCTGAGTGCTGCTCT 96

Search completed: April 25, 2003, 00:00:24
Job time : 56.0118 secs

GenCore version 5.1.4.p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:49:04 ; Search time 373.284 seconds
(without alignments)
4815.901 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtggtatcttcttgcctt.....gcctgagtgcgtcttact 111

Scoring table: OLIGO-MWC
Gapop 60.0 , Gapext 60.0

Searched: 16154066 seqs, 8097743376 residues

Gap size : 0

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estnu:*
5: em_estrov:*
6: em_estrpl:*
7: em_estro:*
8: em_hic:*
9: gb_est1:*
10: gb_est2:*
11: gb_hic:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_iny:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_man:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	470	17	AQ770688
2	45	40.5	525	17	AQ165256 HS_3025_B
3	38	34.2	628	17	AQ237815
4	35	31.5	410	17	AQ442274
5	35	31.5	453	17	AQ437684
6	35	31.5	633	17	AQ390599

c	7	34	30.6	529	17	AQ881246
c	8	33	29.7	482	17	AQ320567
c	9	33	29.7	563	17	AQ420187
c	10	33	29.7	723	17	AQ286439
c	11	31	27.9	553	17	AQ521751
c	12	31	27.9	695	17	AG179297
c	13	30	27.0	360	17	AQ207172
c	14	30	27.0	399	17	AQ115544
c	15	30	27.0	435	17	AQ116061
c	16	30	27.0	551	17	AQ569689
c	17	29	26.1	514	17	AQ003326
c	18	29	26.1	615	17	AG161224
c	19	28	25.2	363	17	AQ120796
c	20	28	25.2	376	17	AQ548294
c	21	28	25.2	401	17	AQ568089
c	22	28	25.2	419	17	AQ362209
c	23	28	25.2	530	17	AQ193128
c	24	28	25.2	553	17	AG160919
c	25	27	24.3	553	17	AQ238365
c	26	27	24.3	606	17	AQ350708
c	27	26	23.4	478	17	B75615
c	28	26	23.4	653	17	AG143347
c	29	26	23.4	690	17	AG11263
c	30	25	22.5	411	17	AQ715895
c	31	25	22.5	427	17	AQ697116
c	32	25	22.5	456	17	AQ707169
c	33	25	22.5	468	17	AQ533287
c	34	25	22.5	519	17	AQ519549
c	35	25	22.5	520	17	AQ375479
c	36	25	22.5	552	17	AQ378145
c	37	25	22.5	563	17	AQ379371
c	38	25	22.5	583	17	AG070759
c	39	25	22.5	609	17	AG155327
c	40	25	22.5	747	17	AG170366
c	41	24	21.6	413	17	AQ266558
c	42	24	21.6	425	17	AQ683450
c	43	24	21.6	471	17	AQ677950
c	44	24	21.6	506	17	AQ284282
c	45	24	21.6	541	17	AQ686074

ALIGNMENTS

RESULT 1
LOCUS AQ770688 470 bp DNA linear GSS 28-JUN-1999
DEFINITION HS_5368_B2_C08-SP66 RPC1-11 Human Male BAC Library Homo sapiens
genomic clone Plate-944 Col-16 Row-F, DNA sequence.

ACCESSION AQ770688
VERSION
KEYWORDS
SOURCE
ORGANISM

human.

Homo sapiens
Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL
MEDLINE
COMMENT

99380589

Contract: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPC1-11. For BAC
library availability, please contact Pieter de Jong

(pieterdejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering/bac.htm>) or from Research Genetics (<http://www.hsc.washington.edu>) or from Research Genetics (<http://inforesgen.com>). BAC end Web Server: <http://www.hsc.washington.edu>

Plate: 944 row: F column: 16

Seq primer: SP6

Class: BAC ends

High quality sequence stop: 470.

FEATURES
Location/Qualifiers

1..470

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate=944 Col=16 Row=F"

/clone_lib="RPCI-11 Human Male BAC Library"

/sex="male"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"

ORIGIN
E COUNT 83 a 112 c 131 g 141 t 3 others

Query Match 100.0%; Score 111; DB 17; Length 470;
Best Local Similarity 100.0%; Pred. No. 2.8e-47;
Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGGGCGATCTTTGCGCTGCGAGATCTTTTATCTTTCAGGAGCTTGGGGCCG 60
|||||
Db 103 ATGGGCGATCTTTGCGCTGCGAGATCTTTTATCTTTCAGGAGCTTGGGGCCG 162

OY 61 GAGTATGTAACCTCTGCGCTGCTGTGCTGCTGAGTGGCTGCTACT 111
|||||
Db 163 GAGTATGTAACCTCTGCGCTGCTGTGTGTGCTGCTGAGTGGCTGCTACT 213

RESULT 2
LOCUS A0165256 525 bp DNA linear GSS 16-OCT-1998

DEFINITION HS.3025.B2.G06.T7 CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone Plate=3025 Col=12 Row=N, DNA sequence.

ACCESSION A0165256
VERSION A0165256.1 GI:3563451
KEYWORDS GSS.

SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 525)
Mahaitas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D., and
Hood,L.

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

TITLE
JOURNAL
MEDLINE
COMMENT
CONTACT: Mahaitas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887

Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3025 row: N column: 12
Class: BAC ends

High quality sequence stop: 525.

FEATURES
Location/Qualifiers

1..525
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3025 Col=12 Row=N"
/clone_lib="CIT Approved Human Genomic Sperm Library D"

/sex="male"
/note="Organ: sperm; Vector: pBelosBAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT 102 a 139 c 137 g 143 t 4 others

Query Match 40.5%; Score 45; DB 17; Length 525;
Best Local Similarity 100.0%; Pred. No. 7.7e-13;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 34 TCATCTTTCGAGGACCTTGGGCGGAGATGATTAACCTCG 78
|||||
Db 99 TCATCTTTCGAGGACCTTGGGCGGAGATGATTAACCTCG 143

RESULT 3
LOCUS A0237815 628 bp DNA linear GSS 21-APR-1999

DEFINITION RPC111-70H4.TK RPCI-11 Homo sapiens genomic clone RPCI-11-70H4, DNA
sequence.

ACCESSION A0237815
VERSION A0237815.1 GI:3670106
KEYWORDS GSS.

SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 628)
Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., White,C., de Jong,P. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)

TITLE
JOURNAL
COMMENT
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208

Email: mdadams@tigr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genetics (<http://inforesgen.com>). BAC end search page:
http://www.tigr.org/vdb/humgen/bac_end_search.html

Seq primer: T7
Class: BAC ends.

FEATURES
Location/Qualifiers

1..628
/organism="Homo sapiens"
/db_xref="taxon:9606"
/db_xref="GDB:752667"

/clone="RPCI-11-70H4"
/clone_lib="RPCI-11"
/sex="male"

/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC111 Human Male BAC Library"

BASE COUNT 125 a 150 c 164 g 189 t

Query Match 34.2%; Score 38; DB 17; Length 628;
Best Local Similarity 100.0%; Pred. No. 3.5e-09;
Matches 38; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 61 GAGTATGTAACCTCTGCGCTGCTGTGCTGCTGAG 98
|||||
Db 157 GAGTATGTAACCTCTGCGCTGCTGTGCTGCTGAG 194

RESULT 4
LOCUS A0442274 410 bp DNA linear GSS 31-MAR-1999

A0442274

DEFINITION HS_5137_A1-F12.SP6E RPCI-11 Human Male BAC Library Homo sapiens
ACCESSION A0442274
VERSION A0442274
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 410)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
or from Research Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: K column: 23
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 410.
FEATURES
Location/Qualifiers
source
1..410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=23 Row=K"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBAC3.6 vector at EcoRI sites"
BASE COUNT 80 a 111 c 111 g 107 t 1 others
ORIGIN
Query Match 31.5%; Score 35; DB 17; Length 410;
Best Local Similarity 100.0%; Pred. No. 1.2e-07;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 TCTGTGGTCTGTGTGTGCTGAGTGGCTGCTCT 108
|||||
Db 160 TCTGTGGTCTGTGTGTGCTGAGTGGCTGCTCT 194
RESULT 5
LOCUS A0437684 453 bp DNA linear GSS 31-MAR-1999
DEFINITION HS_5137_A2.SP6E RPCI-11 Human Male BAC Library Homo sapiens
ACCESSION A0437684
VERSION A0437684
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 453)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
or from Research Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: O column: 12
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 453.
FEATURES
Location/Qualifiers
source
1..453
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=12 Row=O"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBAC3.6 vector at EcoRI sites"
BASE COUNT 84 a 127 c 117 g 124 t 1 others
ORIGIN
Query Match 31.5%; Score 35; DB 17; Length 453;
Best Local Similarity 100.0%; Pred. No. 1.2e-07;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 TCTGTGGTCTGTGTGTGCTGAGTGGCTGCTCT 108
|||||
Db 163 TCTGTGGTCTGTGTGTGCTGAGTGGCTGCTCT 197
RESULT 6
LOCUS A0390599/c 635 bp DNA linear GSS 06-MAR-1999
DEFINITION CITBI-E1-2544B15.TR CITBI-E1 Homo sapiens genomic clone 2544B15,
DNA sequence.
ACCESSION A0390599
VERSION A0390599.1 GI:4361622
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 635)
AUTHORS Zhao,S., Adams,M.D., Nierman,W., Malek,J., Shizuya,H., Simon,M. and
Venter,J.C.
TITLE Use of BAC End Sequences from Caltech Libraries for Sequence-Ready
Map Building
JOURNAL Unpublished (1997)
COMMENT Other GSSs: CITBI-E1-2544B15.TR
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
7712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: bheer@igrg.org
Clones are available from Research Genetics (info@resgen.com). BAC

end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13 Reverse
Class: BAC ends.

FEATURES
Source

Location/Qualifiers
1. 635
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="2544815"
/clone_lib="CITBI-E1"
/sex="male"
/cell_type="sperm"
/note="Vector: pBeloBAC11; Site_1: EcoRI; site_2: EcoRI;
Caltech Human BAC Library D"
BASE COUNT 192 a 144 c 172 g 127 t
ORIGIN

Query Match 31.5%; Score 35; DB 17; Length 635;
Best Local Similarity 100.0%; Pred. No. 1.3e-07;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
74 TCGTGGCTCTGTGTGTGCTGAGTGGCTCT 108
|||||
DB 527 TCGTGGCTCTGTGTGTGCTGAGTGGCTCT 493

RESULT 7
AO881246/c 529 bp DNA linear GSS 09-NOV-1999
LOCUS AO881246
DEFINITION HS_5137_B1_F08_T7 RPTI-11 Human Male BAC Library Homo sapiens
ACCESSION AO881246
VERSION AO881246.1 GI:6312713
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 529)
Maitra, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T.,
Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D., and
Hood, L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589

CONTACT: Maitra, G.G., Wallace, J.C., Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPTI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buflalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buflalo.edu/ordering_bac.htm)
or from Research Genetics (<http://www.resgen.com>). BAC end Web Server:
<http://www.htsc.washington.edu>
Plate: 8905 row: L column: 15
Seq primer: T7
Class: BAC ends
High quality sequence stop: 529.
Location/Qualifiers
1. 529

FEATURES
Source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate:8905 Col-15 Row=L"
/clone_lib="RPTI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and

EcoRI Methylase. Size selected DNA was cloned into the
pBAC3.6 vector at EcoRI sites"
BASE COUNT 137 a 147 c 121 g 117 t 7 others
ORIGIN

Query Match 30.6%; Score 34; DB 17; Length 529;
Best Local Similarity 100.0%; Pred. No. 4.2e-07;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 60 GGAGTATGTAACCTCTGGCTCTGTGTGTC 93
|||||
DB 382 GGAGTATGTAACCTCTGGCTCTGTGTGTC 349

RESULT 8
AO320567/c 482 bp DNA linear GSS 04-MAY-1999
LOCUS AO320567
DEFINITION RPTI11-99N1.TV RPTI-11 Homo sapiens genomic clone RPTI-11-99N1, DNA
sequence.
ACCESSION AO320567
VERSION AO320567.1 GI:4050696
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 482)
Adams, M.D., Rounsley, S.D., Zhao, S., Bass, S., Linher, K., Golden, K.,
Berry, K., Granger, D., Suh, E., Wible, C., de Jong, P., and Venter, J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)
CONTACT: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeet@tigr.org

Clones are derived from the human BAC library RPTI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buflalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buflalo.edu/ordering_bac.htm)
Research Genetics (<http://www.resgen.com>). BAC end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html
Seq primer: T7
Class: BAC ends.

FEATURES
Source

Location/Qualifiers
1. 482
/organism="Homo sapiens"
/db_xref="GDB:7537944"
/db_xref="taxon:9606"
/clone="RPTI-11-99N1"
/clone_lib="RPTI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPTI11 Human Male BAC Library"
BASE COUNT 142 a 121 c 106 g 112 t 1 others
ORIGIN

Query Match 29.7%; Score 33; DB 17; Length 482;
Best Local Similarity 100.0%; Pred. No. 1.4e-06;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 53 TGGGGCCGGAGTATGTAACCTCTGGCTCT 85
|||||
DB 338 TGGGGCCGGAGTATGTAACCTCTGGCTCT 306

RESULT 9
AO420187/c 563 bp DNA linear GSS 23-MAR-1999
LOCUS AO420187
DEFINITION RPTI-11-185J19.TV RPTI-11 Homo sapiens genomic clone RPTI-11-185J19

, DNA sequence.
 ACCESSION AQ420187
 VERSION AQ420187.1 GI:4477911
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 563)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
 ,J.C.
 TITLE Use of BAC end Sequences from Library RPCI-11 for Sequence-Ready
 Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI-11-185J19.TV
 Contact: Shaying Zhao, William Niernan, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genet cs (info@resgen.com). BAC end search page:
 http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 Seq primer: SP6
 Class: BAC ends.
 FEATURES
 source
 Location/Qualifiers
 1..563
 /organism="Homo sapiens"
 /db_xref="GDB:7570890"
 /db_xref="taxon:9606"
 /clone="RPCI-11-185J19"
 /clone_11b="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: pBACe3.6; Site:1: EcoRI; Site:2: EcoRI;
 RPCI11 Human Male BAC Library"
 BASE COUNT 173 a 127 c 114 g 149 t
 ORIGIN
 Query Match 29.7%; Score 33; DB 17; Length 563;
 Best Local Similarity 100.0%; Pred. No. 1.4e-06;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 53 TGGGGCCGAGTATGTAAACCTCGGTCCTC 85
 ||||||||||||||||||||||||||||||||
 333 TGGGGCCGAGTATGTAAACCTCGGTCCTC 301
 RESULT 10
 AO386439/c 733 bp DNA linear GSS 21-MAY-1999
 LOCUS RPCI11-154D6.TV RPCI-11 Homo sapiens genomic clone RPCI-11-154D6,
 DEFINITION DNA sequence.
 ACCESSION AO386439
 VERSION AO386439.1 GI:4357462
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 723)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
 ,J.C.
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
 Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI11-154D6.TV
 Contact: Shaying Zhao, William Niernan, Mark Adams

Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genomics (info@resgen.com). BAC end search page:
 http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html
 Seq primer: SP6
 Class: BAC ends.
 FEATURES
 source
 Location/Qualifiers
 1..723
 /organism="Homo sapiens"
 /db_xref="GDB:7558829"
 /db_xref="taxon:9606"
 /clone="RPCI-11-154D6"
 /clone_11b="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: pBACe3.6; Site:1: EcoRI; Site:2: EcoRI;
 RPCI11 Human Male BAC Library"
 BASE COUNT 224 a 165 c 135 g 199 t
 ORIGIN
 Query Match 29.7%; Score 33; DB 17; Length 723;
 Best Local Similarity 100.0%; Pred. No. 1.5e-06;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 53 TGGGGCCGAGTATGTAAACCTCGGTCCTC 85
 ||||||||||||||||||||||||||||||||
 Db 331 TGGGGCCGAGTATGTAAACCTCGGTCCTC 299
 RESULT 11
 A2521751
 LOCUS RPCI-11-175G22.TVB RPCI-11 Homo sapiens genomic clone
 DEFINITION DNA sequence.
 ACCESSION A2521751
 VERSION A2521751.1 GI:10834261
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 553)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
 ,J.C.
 TITLE BAC end sequences of library RPCI-11
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI-11-175G22.TV
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genet cs (info@resgen.com). BAC end search page:
 http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 This BAC end was generated during the R&D process and may have
 higher chance of clone tracking errors.
 Seq primer: T7
 Class: BAC ends.
 Location/Qualifiers

FEATURES

```

source
1. .553
/organism="Homo sapiens"
/db_xref="GDB:7566981"
/db_xref="taxon:9606"
/clone="RP43-051111.TJ"
/clone_11b="RP43-051111.TJ"
/sex="Male"
/cell_type="Lymphocytes"
/notes="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RP43 Human Male BAC Library"
BASE COUNT      109 a      148 c      137 g      138 t      1 others
ORIGIN
Query Match      27.9%; Score 31; DB 17; Length 553;
Best Local Similarity 100.0%; Pred. No. 1.6e-05;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 68 TAAACTCTGGCTCTGTGTGTGCTGAG 98
|||||
140 TAAACTCTGGCTCTGTGTGTGCTGAG 170

RESULT 12
AG179297      695 bp      DNA      linear      GSS 09-JAN-2002
LOCUS
Pan troglodytes DNA, clone: RP43-051111.TJ, genomic survey
DEFINITION
sequence.
ACCESSION
AG179297
VERSION
AG179297.1 GI:16708977
KEYWORDS
GSS.
Pan troglodytes male lymphocytes DNA, clone_11b:RP43-43 Chimpanzee
Male BAC Library clone:RP43-051111.TJ.
SOURCE
Pan troglodytes
Organism
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
REFERENCE
1 Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
Toloki,Y., Watanabe,H. and Sakaki,Y.
BAC end sequences of Library RP43-43
2 (bases 1 to 695)
Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
Toloki,Y., Watanabe,H. and Sakaki,Y.
Direct Submission
Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC):
1-7-22 Suehiro-chou,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail:chiimpesgsc.riken.go.jp, URL:http://hnp.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the chimpanzee BAC library RP43-43 This BAC
clone was generated during the R&D process and may have higher chance
of clone tracking errors.
PRIMERS
Sequencing: TJ
LIBRARY
Vector : pBAC3.6
R.Site 1 : EcoRI
R.Site 2 : EcoRI.
FEATURES
Location/Qualifiers
1. 695
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="RP43-051111.TJ"
/sex="male"
/cell_type="Lymphocytes"
/clone_11b="RP43-43 Chimpanzee Male BAC Library"
BASE COUNT      243 a      135 c      122 g      192 t      3 others
ORIGIN
Query Match      27.9%; Score 31; DB 17; Length 695;
Best Local Similarity 100.0%; Pred. No. 1.6e-05;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 61 GAGTATGTAAACTCTGGCTCTGTCTGTCT 91
|||||
Db 171 GAGTATGTAAACTCTGGCTCTGTCTGTCT 201

RESULT 13
AO207172      360 bp      DNA      linear      GSS 18-SEP-1998
LOCUS
H5-3239_B1.C03.T7 CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone Plate=3239 Col=5 Row=F, DNA sequence.
DEFINITION
AO207172
ACCESSION
AO207172.1 GI:3618377
VERSION
GSS.
KEYWORDS
human.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 360)
Mahairas,G.C., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,D., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3239 row: F column: 5
Class: BAC ends
High quality sequence stop: 360.
FEATURES
Location/Qualifiers
1. 360
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3239 Col=5 Row=F"
/clone_11b="CIT Approved Human Genomic Sperm Library D"
/sex="male"
/notes="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coli DH10B"
BASE COUNT      63 a      93 c      107 g      93 t      4 others
ORIGIN
Query Match      27.0%; Score 30; DB 17; Length 360;
Best Local Similarity 100.0%; Pred. No. 4.9e-05;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 79 GGTCTGTGTGTGCTGAGTGGCTGCTCT 108
|||||
Db 171 GGTCTGTGTGTGCTGAGTGGCTGCTCT 200

RESULT 14
AO115544/c      399 bp      DNA      linear      GSS 20-APR-1999
LOCUS
RP43-57K21.YK RP43-11 Homo sapiens genomic clone RP43-11-57K21,
DNA sequence.
DEFINITION
AO115544
ACCESSION
AO115544
VERSION
AO115544.1 GI:3491665
KEYWORDS
GSS.
SOURCE
human.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 399)
Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
TITLE

```

JOURNAL

Unpublished (1998)
Other_GSSs: RPC11-57K21.TJ

Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208

Email: mdamas@tigr.org
Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html
Class: BAC ends.

FEATURES

Location/Qualifiers
1..399

/organism="Homo sapiens"
/db_xref="GDB:7521764"
/db_xref="taxon:9606"
/clone="RPC1-11-57K21"
/clone_lib="RPC1-11"
/sex="Male"

/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"

BASE COUNT 120 a 111 c 95 g 73 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 5e-05;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCGTCTGTGTG 90
|||||

Db 380 GAGTATGTAACCTCGTCTGTGTG 351

RESULT 15

AQ116061/c

LOCUS

DEFINITION

Accession

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (1998)

Other_GSSs: RPC11-57L19.TJ

Contact: Mark Adams

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: mdamas@tigr.org

Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html
Class: BAC ends.

Location/Qualifiers

1..435

FEATURES

Location/Qualifiers
1..435

/organism="Homo sapiens"
/db_xref="GDB:7521786"

/db_xref="taxon:9606"

/clone="RPC1-11-57L19"

/clone_lib="RPC1-11"

/sex="Male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"

BASE COUNT 131 a 115 c 103 g 86 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 5e-05;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCGTCTGTGTG 90
|||||

Db 379 GAGTATGTAACCTCGTCTGTGTG 350

Search completed: April 25, 2003, 00:52:48
Job time : 378.284 secs

[illegible]

```

; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,636
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,874
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,910
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,864
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,631
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,845
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,892
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/057,761
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/047,595
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,599
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,588
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,585
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,586
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,590
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,594
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,589
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,593
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,614
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/043,578
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/043,576
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/047,501
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/043,670
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/056,632
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,664
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,876
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,881
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,909
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,875
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,862
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,887
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,908
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/057,650
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 60/056,884
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/057,669
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 60/049,610
; EARLIER FILING DATE: 1997-06-13

```

```

; EARLIER APPLICATION NUMBER: 60/061,060
; EARLIER FILING DATE: 1997-10-02

```

```

Query Match 14.4%; Score 16; DB 4; Length 997;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 8 GATCTTTGCCTTGA 23
|||||
Db 662 GATCTTTGCCTTGA 677

```

```

RESULT 3
US-09-233-506-1
; Sequence 1, Application US/09233506
; Patent No. 6136580

```

```

; GENERAL INFORMATION:

```

```

; APPLICANT: Fukuda, Minoru
; TITLE OF INVENTION: A Beta-1-6-N-Acetylglucosaminyltransferase That Forms
; FILE REFERENCE: P-1j 3415
; CURRENT APPLICATION NUMBER: US/09/233,506

```

```

; CURRENT FILING DATE: 1999-01-19
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: PatentIn Ver. 2.0

```

```

; SEQ ID NO 1
; LENGTH: 2128

```

```

; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:

```

```

; NAME/KEY: CDS
; LOCATION: (354)..(1670)
US-09-233-506-1

```

```

Query Match 14.4%; Score 16; DB 3; Length 2128;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 8 GATCTTTGCCTTGA 23
|||||
Db 1821 GATCTTTGCCTTGA 1836

```

```

RESULT 4
US-09-334-601-6/c
; Sequence 6, Application US/09334601
; Patent No. 6280989

```

```

; GENERAL INFORMATION:
; APPLICANT: Kapitonov, Dmitri

```

```

; TITLE OF INVENTION: NOVEL SLALYTRANSPERASES
; FILE REFERENCE: VCUIP-6
; CURRENT APPLICATION NUMBER: US/09/334,601

```

```

; CURRENT FILING DATE: 1999-06-17
; NUMBER OF SEQ ID NOS: 94
; SOFTWARE: PatentIn Ver. 2.0

```

```

; SEQ ID NO 6
; LENGTH: 2178
; TYPE: DNA
; ORGANISM: Homo sapiens

```

```

US-09-334-601-6
Query Match 14.4%; Score 16; DB 4; Length 2178;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 26 ATCTTTTCATCTTT 41
|||||
Db 2150 ATCTTTTCATCTTT 2135

```

```

RESULT 5

```

US-09-334-601-1/C
; Sequence 1, Application US/09334601
; Patent No. 6280989
; GENERAL INFORMATION:
; APPLICANT: Kapitonov, Dmitri
; APPLICANT: Yu, Robert
; TITLE OF INVENTION: NOVEL STALYLTRANSFERASES
; FILE REFERENCE: VCUJP-6
; CURRENT APPLICATION NUMBER: US/09/334,601
; CURRENT FILING DATE: 1999-06-17
; NUMBER OF SEQ ID NOS: 94
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 2288
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (29)..(1282)
09-334-601-1

Query Match 14.4%; Score 16; DB 4; Length 2288;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ATCTTTTCATCTTT 41
|||||
DB 2260 ATCTTTTCATCTTT 2245

RESULT 6
US-07-746-705A-16
; Sequence 16, Application US/07746705A
; Patent No. 5451516
; GENERAL INFORMATION:
; APPLICANT: Matthews, Benjamin F.
; APPLICANT: Weisemann, Jane M.
; TITLE OF INVENTION: A Recombinant DNA Molecule Encoding
; TITLE OF INVENTION: a Bifunctional Plant Enzyme: Aspartokinase and Homoserine
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle S. Graeter
; STREET: Bldg. 005, Room 402, BARC-W
; CITY: Beltsville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20705

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/746,705A
FILING DATE: 19910816
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Graeter, Janelle S.
REGISTRATION NUMBER: 35,024
REFERENCE/DOCKET NUMBER: 4000.91
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301)504-5676
TELEFAX: (301)504-5060
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 2915 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: both
TOPOLOGY: linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHETICAL: NO
ANTI-SENSE: NO

FEATURE:
; NAME/KEY: CDS
; LOCATION: 2..2593
US-07-746-705A-16

Query Match 14.4%; Score 16; DB 1; Length 2915;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 CTTTGACGAGCTTCT 53
|||||
DB 1977 CTTTGACGAGCTTCT 1992

RESULT 7
US-08-380-182-18
; Sequence 18, Application US/08380182
; Patent No. 5858749
; GENERAL INFORMATION:
; APPLICANT: Matthews, Benjamin F.
; APPLICANT: Weisemann, Jane M.
; TITLE OF INVENTION: A Bifunctional Protein From Carrots
; TITLE OF INVENTION: (Daucus carota) with Aspartokinase and Homoserine
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle S. Graeter
; STREET: Room 411, Bldg. 005, BARC-W
; CITY: Beltsville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20705

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/380,182
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Graeter, Janelle S.
REGISTRATION NUMBER: 35,024
REFERENCE/DOCKET NUMBER: 0226.94
TELECOMMUNICATION INFORMATION:
TELEPHONE: 301-504-6629
TELEFAX: 301-504-5060
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 2915 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Daucus carota
FEATURE:
NAME/KEY: CDS
LOCATION: 2..2593
US-08-380-182-18

Query Match 14.4%; Score 16; DB 2; Length 2915;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 CTTTGACGAGCTTCT 53
|||||
DB 1977 CTTTGACGAGCTTCT 1992

RESULT 8
US-09-334-601-5/c
Sequence 5, Application US/09334601
Patent No. 6280989
GENERAL INFORMATION:
APPLICANT: Kapitonov, Dmitri
APPLICANT: Yu, Robert
TITLE OF INVENTION: NOVEL STALYLTRANSFERASES
FILE REFERENCE: YCUIP-6
CURRENT APPLICATION NUMBER: US/09/334,601
CURRENT FILING DATE: 1999-06-17
NUMBER OF SEQ ID NOS: 94
SOFTWARE: Patentln Ver. 2.0
SEQ ID NO 5
LENGTH: 3494
TYPE: DNA
ORGANISM: Homo sapiens
US-09-334-601-5

Query Match 14.4%; Score 16; DB 4; Length 3494;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ATCTTTTCATCTTT 41
|||||
DB 3467 ATCTTTTCATCTTT 3452

RESULT 9
US-09-024-020B-1/c
Sequence 1, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLYVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION NUMBER:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5977 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)
US-09-024-020B-1

Query Match 14.4%; Score 16; DB 3; Length 5977;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 80 GTCTGTGTGTGCTT 95
|||||
DB 5814 GTCTGTGTGTGCTT 5799

RESULT 10
US-09-425-043-1/c
Sequence 1, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLYVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5977 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-425-043-1

Query Match 14.4%; Score 16; DB 4; Length 5977;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 80 GTCTGTGTGTGCTT 95
|||||
DB 5814 GTCTGTGTGTGCTT 5799

RESULT 11
US-09-024-020B-2/c

Sequence 2, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 6007 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-2

Query Match
Best Local Similarity 14.4%; Score 16; DB 3; Length 6007;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 80 GTCTCTGTGTGCTCT 95
5844 GTCTCTGTGTGCTCT 5829

RESULT 12
US-09-425-043-2/c
Sequence 2, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA

COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 6007 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-425-043-2

Query Match
Best Local Similarity 14.4%; Score 16; DB 4; Length 6007;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 80 GTCTCTGTGTGCTCT 95
5844 GTCTCTGTGTGCTCT 5829

RESULT 13
US-09-024-020B-7/c
Sequence 7, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997

ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 6556 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-7

Query Match 14.4%; Score 16; DB 3; Length 6556;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

80 GTCTCTGTGTGTCCT 95
|||||
DB 5961 GTCTCTGTGTGTCCT 5946

RESULT 14
US-09-425-043-7/c
Sequence 7, Application US/09425043
Patent No. 6335172

GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 6556 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)

US-09-425-043-7

Query Match 14.4%; Score 16; DB 4; Length 6556;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 GTCTCTGTGTGTCCT 95
|||||
DB 5961 GTCTCTGTGTGTCCT 5946

RESULT 15
US-09-024-020B-43/c
Sequence 43, Application US/09024020B
Patent No. 6030810

GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 43:
SEQUENCE CHARACTERISTICS:
LENGTH: 6586 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-43

Query Match 14.4%; Score 16; DB 3; Length 6586;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 GTCTCTGTGTGTCCT 95
|||||
DB 5991 GTCTCTGTGTGTCCT 5976

Search completed: April 25, 2003, 00:54:06
Job time : 19.1361 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:41:49 ; Search time 22.7692 Seconds

(without alignments)
5304.628 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgagcttcttgcctt.....gccttgatgctgctctact 111

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 709820 seqs, 544064369 residues

Database size: 0
Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

Published Applications_NA: *
1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq: *
2: /cgn2_6/ptodata/1/pubpna/PCOT_NEW_PUB.seq: *
3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq: *
4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq: *
5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq: *
6: /cgn2_6/ptodata/1/pubpna/PCOTUS_PUBCOMB.seq: *
7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq: *
8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq: *
9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq: *
10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq: *
11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq: *
12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq: *
13: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq: *
14: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	33	29.7	664973	10 US-09-263-959-1	Sequence 1, Appli
2	18	16.2	431	10 US-09-866-562-80	Sequence 80, Appl
3	18	16.2	431	10 US-09-866-562-87	Sequence 87, Appl
4	18	16.2	570	10 US-09-864-761-9118	Sequence 9118, Ap
5	17	15.3	531	9 US-10-692-154-1899	Sequence 1899, Ap
6	17	15.3	531	10 US-09-764-847-1899	Sequence 1899, Ap
7	17	15.3	609	10 US-09-874-300-8084	Sequence 8084, Ap
8	17	15.3	1047	10 US-09-822-830A-468	Sequence 468, Appl
9	17	15.3	172637	10 US-09-805-458A-3	Sequence 3, Appli
10	17	15.3	1503841	9 US-09-946-807-1	Sequence 1, Appli
11	17	15.3	1503841	10 US-09-795-668-1	Sequence 1, Appli
12	17	15.3	1503841	10 US-09-795-668-1	Sequence 1, Appli
13	16	14.4	281	10 US-09-964-824A-385	Sequence 385, App
14	16	14.4	374	9 US-10-046-935-2033	Sequence 2033, Ap
15	16	14.4	374	9 US-09-878-178-2033	Sequence 2033, Ap
16	16	14.4	374	9 US-10-146-502-2033	Sequence 2033, Ap
17	16	14.4	374	9 US-10-060-036-985	Sequence 985, App
18	16	14.4	374	9 US-10-060-036-2528	Sequence 2528, Ap
19	16	14.4	436	9 US-09-918-995-26108	Sequence 26108, A

C 20	16	14.4	447	10 US-09-880-107-871	Sequence 871, App
C 21	16	14.4	481	10 US-09-560-863-45	Sequence 45, Appl
C 22	16	14.4	494	9 US-09-918-995-20796	Sequence 20796, A
C 23	16	14.4	497	10 US-09-864-761-1356	Sequence 1356, Ap
C 24	16	14.4	524	10 US-09-797-207-5	Sequence 5, Appli
C 25	16	14.4	548	10 US-09-864-761-12421	Sequence 12421, A
C 26	16	14.4	552	10 US-09-998-598-1325	Sequence 1325, Ap
C 27	16	14.4	557	10 US-09-764-877-746	Sequence 746, App
C 28	16	14.4	557	10 US-09-764-877-3495	Sequence 3495, Ap
C 29	16	14.4	747	12 US-10-001-879-101	Sequence 101, App
C 30	16	14.4	805	9 US-09-984-245-106	Sequence 106, App
C 31	16	14.4	805	9 US-09-966-262-106	Sequence 106, App
C 32	16	14.4	805	9 US-09-983-966-106	Sequence 106, App
C 33	16	14.4	805	9 US-10-143-090-106	Sequence 106, App
C 34	16	14.4	997	9 US-09-809-391-307	Sequence 307, App
C 35	16	14.4	1009	10 US-09-764-864-12	Sequence 12, Appl
C 36	16	14.4	2002	10 US-09-925-300-592	Sequence 592, App
C 37	16	14.4	2017	9 US-10-102-806-137	Sequence 137, App
C 38	16	14.4	2108	10 US-09-797-207-3	Sequence 3, Appli
C 39	16	14.4	2147	9 US-09-981-353-43	Sequence 43, Appl
C 40	16	14.4	2229	10 US-09-925-297-337	Sequence 337, App
C 41	16	14.4	2268	9 US-10-102-806-142	Sequence 142, App
C 42	16	14.4	2287	10 US-09-764-864-473	Sequence 473, App
C 43	16	14.4	2319	10 US-09-874-390-1	Sequence 1, Appli
C 44	16	14.4	2340	9 US-09-964-245-51	Sequence 51, Appl
C 45	16	14.4	2340	9 US-09-966-262-51	Sequence 51, Appl

ALIGNMENTS

RESULT 1
US-09-263-959-1/c
Sequence 1, Application US/09263959
Patent No. US20020150891A1
GENERAL INFORMATION:
APPLICANT: Hood, Leroy E.
APPLICANT: Rowen, Lee
TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U
NUMBER OF SEQUENCES: 1279
CORRESPONDENCE ADDRESS:
ADDRESSEE: Seed and Berry LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: US
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/263,959
FILING DATE: 05-MAR-1999
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 664973 base pairs
TYPE: nucleic acid
STRANDEDNESS: Single
TOPOLOGY: linear
US-09-263-959-1
Query Match 29.7%; Score 33; DB 10; Length 664973;

Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 66 TGTAAACTCTGGCTCTGTGTGTGCTGAG 98
|||||

Db 404690 TGTAAACTCTGGCTCTGTGTGTGCTGAG 404658

RESULT 2

US-09-866-562-80
; Sequence 80, Application US/09866562
; Patent No. US2002009758A1
; GENERAL INFORMATION:
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Bangur, Tonglong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Klee, Jennifer
; APPLICANT: Switzer, Anne
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; FILE REFERENCE: 210121.502
; CURRENT APPLICATION NUMBER: US/09/866,562
; CURRENT FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 96
; SEQ ID NO 80
; LENGTH: 431
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 361..431
; OTHER INFORMATION: n = A,T,C or G
US-09-866-562-80

Query Match 16.2%; Score 18; DB 10; Length 431;
Best Local Similarity 100.0%; Pred. No. 3;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 91 TGCCTGAGTGGCTGCTCT 108
|||||

Db 18 TGCCTGAGTGGCTGCTCT 35

RESULT 3

US-09-866-562-87
; Sequence 87, Application US/09866562
; Patent No. US2002009758A1
; GENERAL INFORMATION:
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Bangur, Tonglong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Klee, Jennifer
; APPLICANT: Switzer, Anne
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; FILE REFERENCE: 210121.502
; CURRENT APPLICATION NUMBER: US/09/866,562
; CURRENT FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 96
; SEQ ID NO 87
; LENGTH: 431
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 361..431
; OTHER INFORMATION: n = A,T,C or G
US-09-866-562-87

Query Match 16.2%; Score 18; DB 10; Length 431;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 91 TGCCTGAGTGGCTGCTCT 108
|||||

Db 18 TGCCTGAGTGGCTGCTCT 35

RESULT 4

US-09-864-761-9118/c
; Sequence 9118, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aeomica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263,6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 9118
; LENGTH: 570
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000053.1
; OTHER INFORMATION: EXPRESSED IN HEPA, SIGNAL = 2.8
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
US-09-864-761-9118

Query Match 16.2%; Score 18; DB 10; Length 570;
 Best Local Similarity 100.0%; Pred. No. 3;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 85 TGTGTGCTGCTGAGTGGC 102
 ||||||||||||||||
 Db 505 TGTGTGCTGCTGAGTGGC 488

RESULT 5
 US-10-092-154-1899/c
 ; Sequence 1899, Application US/10092154
 ; Publication No. US200300543755a1
 ; GENERAL INFORMATION:
 ; APPLICANT: Rosen et al.
 ; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
 ; FILE REFERENCE: PC009C1
 ; CURRENT APPLICATION NUMBER: US/10/092,154
 ; CURRENT FILING DATE: 2002-03-07
 ; NUMBER OF SEQ ID NOS: 2003
 ; Prior Application removed - See File Wrapper or Palm
 ; SOFTWARE: PatentIn Ver. 2.0
 ; SEQ ID NO: 1899
 ; LENGTH: 531
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 US-10-092-154-1899

Query Match 15.3%; Score 17; DB 9; Length 531;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
 ||||||||||||||||
 Db 141 TCTTTTCATCTTTGCA 125

RESULT 6
 US-09-764-847-1899/c
 ; Sequence 1899, Application US/09764847
 ; Patent No. US20020132767A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Rosen et al.
 ; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
 ; FILE REFERENCE: PC009
 ; CURRENT APPLICATION NUMBER: US/09/764,847
 ; CURRENT FILING DATE: 2001-01-17
 ; Prior application data removed - consult PALM or file wrapper
 ; NUMBER OF SEQ ID NOS: 2003
 ; SOFTWARE: PatentIn Ver. 2.0
 ; SEQ ID NO: 1899
 ; LENGTH: 531
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 US-09-764-847-1899

Query Match 15.3%; Score 17; DB 10; Length 531;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
 ||||||||||||||||
 Db 141 TCTTTTCATCTTTGCA 125

RESULT 7
 US-09-974-300-8084/c
 ; Sequence 8084, Application US/09974300
 ; Patent No. US20020146721A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Berka, Randy M.
 ; APPLICANT: Clausen, Ib Groth
 ; TITLE OF INVENTION: Methods For Monitoring Multiple Gene

; TITLE OF INVENTION: Expression
 ; FILE REFERENCE: 10085,500-US
 ; CURRENT APPLICATION NUMBER: US/09/974,300
 ; CURRENT FILING DATE: 2001-10-05
 ; PRIOR APPLICATION NUMBER: 09/680,598
 ; PRIOR FILING DATE: 2000-10-06
 ; PRIOR APPLICATION NUMBER: 60/279,526
 ; PRIOR FILING DATE: 2001-03-27
 ; NUMBER OF SEQ ID NOS: 8481
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO: 8084
 ; LENGTH: 609
 ; TYPE: DNA
 ; ORGANISM: Bacillus clausii
 US-09-974-300-8084

Query Match 15.3%; Score 17; DB 10; Length 609;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
 ||||||||||||||||
 Db 536 TCTTTTCATCTTTGCA 520

RESULT 8
 US-09-822-830A-468
 ; Sequence 468, Application US/09822830A
 ; Patent No. US20020142952A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Genetics Institute, Inc.
 ; APPLICANT: Wong, Gordon G.
 ; APPLICANT: Clark, Hilary
 ; APPLICANT: Fecthel, Kim
 ; APPLICANT: Agostino, Michael J.
 ; APPLICANT: Howes, Steven H.
 ; APPLICANT: Resnick, Richard J.
 ; APPLICANT: Gulukota, Kamalakara
 ; APPLICANT: Graham, James R.
 ; TITLE OF INVENTION: POLYNUCLEOTIDES ENCODING NOVEL SECRETED PROTEINS
 ; FILE REFERENCE: GIN 6402
 ; CURRENT APPLICATION NUMBER: US/09/822,830A
 ; CURRENT FILING DATE: 2001-03-29
 ; PRIOR APPLICATION NUMBER: 60/195,604
 ; PRIOR FILING DATE: 2000-04-06
 ; NUMBER OF SEQ ID NOS: 631
 ; SOFTWARE: PatentIn Ver. 2.0
 ; SEQ ID NO: 468
 ; LENGTH: 1047
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 US-09-822-830A-468

Query Match 15.3%; Score 17; DB 10; Length 1047;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 76 CTGGGTCCTGTGTG 92
 ||||||||||||||||
 Db 828 CTGGGTCCTGTGTG 844

RESULT 9
 US-09-805-458A-3
 ; Sequence 3, Application US/09805458A
 ; Patent No. US20020042100A1
 ; GENERAL INFORMATION:
 ; APPLICANT: YAN, Chunhua et al
 ; TITLE OF INVENTION: ISOLATED HUMAN ION CHANNEL PROTEINS,
 ; NUCLEIC ACID MOLECULES ENCODING HUMAN ION CHANNEL PROTEINS,
 ; TITLE OF INVENTION: AND USBS THEREOF
 ; FILE REFERENCE: CI000722
 ; CURRENT APPLICATION NUMBER: US/09/805,458A

```

: CURRENT FILING DATE: 2001-03-14
: NUMBER OF SEQ ID NOS: 6
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 3
: LENGTH: 172637
: TYPE: DNA
: ORGANISM: Human
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(172637)
: OTHER INFORMATION: n = A,T,C or G
: US-09-805-458A-3
```

```

Query Match          15.3%; Score 17; DB 10; Length 172637;
Best Local Similarity 100.0%; Pred. No. 9,7;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```

QY 19 TTGCAGATTCTTTTTC 35
      |||
150045 TTGCAGATTCTTTTTC 150061
```

```

RESULT 10
: US-09-946-807-1
: Sequence 1, Application US/09946807
: Patent No. US20020165144A1
: GENERAL INFORMATION:
: APPLICANT: Stefansson, Hreinn
: APPLICANT: Steinthorsson, Valgerdur
: APPLICANT: Gulcher, Jeffrey R.
: TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
: FILE REFERENCE: 2345,2004-001
: CURRENT APPLICATION NUMBER: US/09/946,807
: CURRENT FILING DATE: 2001-09-05
: PRIOR APPLICATION NUMBER: US/09/795,668
: PRIOR FILING DATE: 2001-02-28
: PRIOR APPLICATION NUMBER: US 09/515,716
: PRIOR FILING DATE: 2000-02-28
: NUMBER OF SEQ ID NOS: 1531
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 1
: LENGTH: 1503841
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: y=t/u or c
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: m=a or c
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: k=g or t/u
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: s=g or c
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: w=a or t/u
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: b=g or c or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: d=a or g or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
```

```

: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: d=a or g or t/u
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: h=a or c or t/u
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: v=a or g or c
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: n=a or g or c or t/u
: US-09-946-807-1
```

```

Query Match          15.3%; Score 17; DB 9; Length 1503841;
Best Local Similarity 100.0%; Pred. No. 9,4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```

QY 26 ATTCTTTTCATCTTGG 42
      |||
Db 268726 ATTCTTTTCATCTTGG 268742
```

```

RESULT 11
: US-09-795-668-1
: Sequence 1, Application US/09795668
: Patent No. US20020045577A1
: GENERAL INFORMATION:
: APPLICANT: Stefansson, Hreinn
: APPLICANT: Steinthorsson, Valgerdur
: APPLICANT: Gulcher, Jeffrey R.
: TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
: FILE REFERENCE: 2345,2004-001
: CURRENT APPLICATION NUMBER: US/09/795,668
: CURRENT FILING DATE: 2001-02-28
: PRIOR APPLICATION NUMBER: US 09/515,716
: PRIOR FILING DATE: 2000-02-28
: NUMBER OF SEQ ID NOS: 1531
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 1
: LENGTH: 1503841
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: r=g or a
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: y=t/u or c
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: m=a or c
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: k=g or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: s=g or c
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: w=a or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: b=g or c or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
: OTHER INFORMATION: d=a or g or t/u
: NAME/KEY: misc_feature
: LOCATION: (1)...(1531)
```



```

OTHER INFORMATION: h=a or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: v=a or g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: n=a or g or c or t/u
US-09-795-668-1

```

```

Query Match
Best Local Similarity 15.3%; Score 17; DB 10; Length 1503841;
Pred. No. 9.4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTTG 42
|||||
Db 268726 ATTCTTTTCATCTTTG 268742

```

```

RESULT 12
US-09-795-666-1
Sequence 1, Application US/09795666
Patent No. US20020094954A1
GENERAL INFORMATION:
APPLICANT: Steinhilber, Hrein
APPLICANT: Steinhilber, Hrein
APPLICANT: Gulcher, Jeffrey R.
TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
FILE REFERENCE: 2345, 2005-001
CURRENT APPLICATION NUMBER: US/09/795,666
CURRENT FILING DATE: 2001-02-28
PRIOR APPLICATION NUMBER: US 09/515,715
PRIOR FILING DATE: 2000-02-28
NUMBER OF SEQ ID NOS: 1531
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 1
LENGTH: 1503841
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: r=g or a
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: y=l/u or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: m=a or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: k=g or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: s=g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: w=a or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: b=g or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: d=a or g or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: h=a or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: v=a or g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: n=a or g or c or t/u
US-09-795-666-1

```

```

Query Match
Best Local Similarity 15.3%; Score 17; DB 10; Length 1503841;
Pred. No. 9.4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTTG 42
|||||
Db 268726 ATTCTTTTCATCTTTG 268742

```

```

RESULT 13
US-09-964-824A-385
Sequence 385, Application US/09964824A
Patent No. US20020102531A1
GENERAL INFORMATION:
APPLICANT: Horigan, Stephen
TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using Sign
FILE REFERENCE: 689290-73
CURRENT APPLICATION NUMBER: US/09/964,824A
CURRENT FILING DATE: 2001-09-27
PRIOR APPLICATION NUMBER: US/60/236,033
PRIOR FILING DATE: 2000-09-28
PRIOR APPLICATION NUMBER: US/60/236,032
PRIOR FILING DATE: 2000-09-28
PRIOR APPLICATION NUMBER: US/60/236,028
PRIOR FILING DATE: 2000-09-28
NUMBER OF SEQ ID NOS: 583
SOFTWARE: Patentin Version 3.0
SEQ ID NO 385
LENGTH: 281
TYPE: DNA
ORGANISM: Homo sapiens
US-09-964-824A-385

```

```

Query Match
Best Local Similarity 14.4%; Score 16; DB 10; Length 281;
Pred. No. 36;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTT 41
|||||
Db 22 ATTCTTTTCATCTTT 37

```

```

RESULT 14
US-10-046-935-2033
Sequence 2033, Application US/10046935
Patent No. US20020156011A1
GENERAL INFORMATION:
APPLICANT: Jiang, Yugu
APPLICANT: Harlocker, Susan L.
APPLICANT: Secret, Heather
APPLICANT: Wang, Aijun
APPLICANT: Stolk, John A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
FILE REFERENCE: 210121.527C1
CURRENT APPLICATION NUMBER: US/10/046,935
CURRENT FILING DATE: 2002-01-15
NUMBER OF SEQ ID NOS: 2239
SOFTWARE: FASTSEQ for Windows Version 4.0
SEQ ID NO 2033
LENGTH: 374
TYPE: DNA
ORGANISM: Homo sapiens
US-10-046-935-2033

```

```

Query Match
Best Local Similarity 14.4%; Score 16; DB 9; Length 374;
Pred. No. 36;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 8 GATCTTTTCCTTGCA 23
|||||

```

Db 108 GATCTTTGCTTGA 123

RESULT 15

US-09-878-178-2033
 ; Sequence 2033, Application US/09878178
 ; Patent No. US20020177552A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Jiang, Yugu
 ; APPLICANT: Harlocker, Susan L.
 ; APPLICANT: Secrist, Heather
 ; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
 ; TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
 ; FILE REFERENCE: 210121.527
 ; CURRENT APPLICATION NUMBER: US/09/878.178
 ; CURRENT FILING DATE: 2001-06-08
 ; NUMBER OF SEQ ID NOS: 2237
 ; SOFTWARE: fastseq for windows version 4.0
 ; SEQ ID NO 2033
 ; LENGTH: 374
 ; TYPE: DNA
 ; ORGANISM: Homo sapien
 US-09-878-178-2033

Query Match 14.4%; Score 16; DB 9; Length 374;
 Best Local Similarity 100.0%; Pred. No. 36;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 GATCTTTGCTTGA 23
 |||||
 Db 108 GATCTTTGCTTGA 123

Search completed: April 25, 2003, 02:09:14
 Job time : 1043.77 secs

GenCore version 5.1.4.p5_4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:20:48 ; Search time 1096.62 seconds
(without alignments)
10509.347 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_446

Perfect score: 396
Sequence: 1 atgggtgacatttgcctt.....gamctgatatcattgtga 396

Scoring table:
OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 2054640 seqs, 14551402878 residues

Size: 0

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: GenEmbl:

1: gb_da:*
2: gb_htg:*
3: gb_in:*
4: gb_cm:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_da:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_on:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_to:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pln:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	299	75.5	169620	2 AC012674	AC012674 Homo sapi
C 2	48	12.1	143372	9 AL137847	AL137847 Human DNA
C 3	40	10.1	123779	30 AC0221025	AC0221025 Homo sapi
C 4	40	10.1	128118	2 AC076969	AC076969 Homo sapi
C 5	40	10.1	128583	9 AC121249	AC121249 Homo sapi
C 6	40	10.1	148290	9 AC107979	AC107979 Homo sapi
C 7	40	10.1	165649	9 AC103996	AC103996 Homo sapi
C 8	40	10.1	178650	9 AC104303	AC104303 Homo sapi
C 9	40	10.1	192826	9 AC090762	AC090762 Homo sapi
C 10	39	9.8	32918	2 AC007445	AC007445 Homo sapi
C 11	39	9.8	38936	9 AL358817	AL358817 Human DNA
C 12	39	9.8	124271	2 AC025179	AC025179 Homo sapi
C 13	39	9.8	146671	2 AC008814	AC008814 Homo sapi
C 14	39	9.8	159747	2 AP001019	AP001019 Homo sapi
C 15	39	9.8	162740	2 AC034249	AC034249 Homo sapi
C 16	39	9.8	169772	9 AC069538	AC069538 Homo sapi
C 17	39	9.8	175466	9 AL607077	AL607077 Human DNA
C 18	38	9.6	110000	2 AC068875	AC068875 Homo sapi
C 19	38	9.6	166706	9 AC068875	AC068875 Homo sapi
C 20	38	9.6	207408	2 AC068618	AC068618 Homo sapi
C 21	38	9.6	207548	9 AC087283	AC087283 Homo sapi
C 22	37	9.3	135090	9 HS164112	AL009028 Homo sapi
C 23	36	9.1	121720	9 AL581491	AL581491 Human DNA
C 24	35	8.8	153940	9 AC022294	AC022294 Homo sapi
C 25	35	8.8	176253	2 AP001078	AP001078 Homo sapi
C 26	35	8.8	205639	2 AP001793	AP001793 Homo sapi
C 27	35	8.8	210734	2 AC015676	AC015676 Homo sapi
C 28	35	8.8	212055	2 AP000899	AP000899 Homo sapi
C 29	35	8.8	325069	2 AC079737	AC079737 Homo sapi
C 30	34	8.6	99577	9 AC026324	AC026324 Homo sapi
C 31	34	8.6	108040	2 AC068150	AC068150 Homo sapi
C 32	34	8.6	134760	9 AC099484	AC099484 Homo sapi
C 33	34	8.6	146059	2 AC019030	AC019030 Homo sapi
C 34	34	8.6	171347	9 AC099776	AC099776 Homo sapi
C 35	34	8.6	172206	9 AC092119	AC092119 Homo sapi
C 36	34	8.6	172567	2 AC015493	AC015493 Homo sapi
C 37	34	8.6	173166	9 AC092375	AC092375 Homo sapi
C 38	34	8.6	177447	2 AC104687	AC104687 Homo sapi
C 39	34	8.6	182881	9 AC090980	AC090980 Homo sapi
C 40	34	8.6	183016	2 AC068611	AC068611 Homo sapi
C 41	34	8.6	193929	9 AC090797	AC090797 Homo sapi
C 42	34	8.6	273807	2 AC025421	AC025421 Homo sapi
C 43	34	8.6	316296	2 AC092285	AC092285 Homo sapi
C 44	33	8.3	33458	9 HSDJ60101	AL109656 Human DNA
C 45	33	8.3	49616	9 AL365267	AL365267 Human DNA

ALIGNMENTS

RESULT 1
AC012674/c 169620 bp DNA linear HTG 07-SEP-2000
LOCUS Homo sapiens chromosome 3 clone RP1-458H3, WORKING DRAFT SEQUENCE,
DEFINITION 18 unordered pieces.
AC012674
AC012674.10 GI:9719580
VERSION HTG: HTGS PHASE1, HTGS_DRAFT.
KEYWORDS
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE
1 (bases 1 to 169620)
Muzny,D.M., Adams,C., Bailey,M., Barbara,J., Blankenburg,K.,
Bodots,B., Bouck,J., Bowie,S., Brooks,A., Bunay,C., Bunac,C.,

TITLE JOURNAL REFERENCE AUTHORS TITLE JOURNAL

COMMENT

Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N., Dugan-Rocha, S., Durbin, K. J., Fernandez, C., Ferraguto, D., Forcman-Tenney, J., Frantz, P., Ganesh, R., Gorrell, J. H., Gorrell, L. L., Guevara, W., Harris, K., Hernandez, J., Hodgson, A., Hogues, M., Holloway, C., Hosak, H., Jackson, L. E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondejewski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Lichtarge, O., Liu, J., Liu, W., Logan, O., Lozano, R. J., Lu, J., Lucier, R., Martin, R., Martinez, C., McLeod, M. P., Mei, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Oswal, G., Parish, B., Paxton, S., Payton, B., Perez, L., Pi, L. L., Quiles, M., Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S., Shah, E., Shen, H., Simon, M., Sparks, A., Stamps, A., Suckang, R., Tabor, P., Taylor, T., Vasquez, L., Vinson, R., Vo, Q., Wahab, M., Wellington, S., Weinstein, G., Weinstein, I. R., Williamson, A., Worley, K., Wren, J., Wrenford, G., Yu, W., Zhou, X., Nelson, D. and Gbbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.
Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu

Project Information
Center project name: HMOG
Center clone name: RP1-458H3

Summary Statistics
Assembly program: Phrap, version 0.990329
Consensus quality: 159025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720: sum-of-coverage estimation
Estimated insert size: 171608: agarose-fp estimation
Quality coverage: 3.9x in Q20 bases; agarose-fp estimation
Quality coverage: 4.1x in Q20 bases; sum-of-coverage estimation

NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 28689: contig of 28689 bp in length
28690 28789: gap of unknown length
28790 50832: contig of 22043 bp in length
50833 50932: gap of unknown length
50933 69144: contig of 18212 bp in length
69145 69244: gap of unknown length
69245 84204: contig of 14960 bp in length
84205 84304: gap of unknown length
84305 94667: contig of 10363 bp in length
94668 94767: gap of unknown length
94768 107261: contig of 12494 bp in length
107262 107361: gap of unknown length
107362 117550: contig of 10189 bp in length
117551 117650: gap of unknown length
117651 126939: contig of 9289 bp in length
126940 127039: gap of unknown length
127040 135040: contig of 8001 bp in length
135041 135140: gap of unknown length
135141 141639: contig of 6499 bp in length

FEATURES
source
1..169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="13"
/clone="RP1-458H3"
BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others
ORIGIN
Query Match 75.5% Score 299; DB 2; Length 169620;
Best Local Similarity 99.7%; Pred. No. 3.2e-170;
Matches 349; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
1 ANGGGATGATCTTTTGGCTTCGAGGATCTTTTCATCTTTCAGAGCACTTCGGGCG 60
|||
DB 87441 ANGGGATGATCTTTTGGCTTCGAGGATCTTTTCATCTTTCAGAGCACTTCGGGCG 87382
QY 61 GAGTATGTAACCTCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTACAGACTCTG 120
|||
DB 87381 GAGTATGTAACCTCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTACAGACTCTG 87322
QY 121 CATACACAGCTCTGTATATTCGAGCCAGGCCCTGTCGATGGCTACAGAGATTC 180
|||
DB 87321 CATACACAGCTCTGTATATTCGAGCCAGGCCCTGTCGATGGCTACAGAGATTC 87262
QY 181 CCTGATCTGTGGGTCGAAAGATCTGTGGAGAACTGTGTTTCTCGATGGGTCAC 240
|||
DB 87261 CCTGATCTGTGGGTCGAAAGATCTGTGGAGAACTGTGTTTCTCGATGGGTCAC 87202
QY 241 AATCCTGCTGCTCCCTGCTGGAGTGGGCTTTTTCGTCATGCTGCTGCC 300
|||
DB 87201 AATCCTGCTGCTCCCTGCTGGAGTGGGCTTTTTCGTCATGCTGCTGCC 87142
QY 301 TGGGGGGGGGATGGCCCTACCCCATTTTTCATCTCTGTGGTCAG 350
|||
DB 87141 TGGGGGGGGGATGGCCCTACCCCATTTTTCATCTCTGTGGTCAG 87092

RESULT 2
AL137847/c 143372 bp DNA linear PRI 16-NOV-2001
LOCUS AL137847 Human DNA sequence from clone RP11-439K3 on chromosome 9q22.2-31.1,
DEFINITION complete sequence.
ACCESSION AL137847
VERSION AL137847.12 GI:16973786
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 143372)
AUTHORS Kimberley, A.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk
On Nov 17, 2001 this sequence version replaced gi:16408610.


```

CC * 120299 122220: contig of 1922 bp in length
CC * 122321 122320: gap of unknown length.
CC * 122321 123779: contig of 1459 bp in length.
XX
FH Key Location/Qualifiers
FH source 1. 123779
FT /chromosome="3"
FT /db_xref="taxon:9606"
FT /organism="Homo sapiens"
FT /clone="RP11-79K17"
XX
SQ Sequence 123779 BP; 37802 A; 24017 C; 23437 G; 37612 T; 911 other:

Query Match 10.1%; Score 40; DB 30; Length 123779;
Best Local Similarity 100.0%; Pred. No. 2, 1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 69 AAAACTCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 108
3083 AAAACTCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 3044

RESULT 4
LOCUS AC076969 128118 bp DNA linear HTG 15-OCT-2001
DEFINITION Homo sapiens chromosome 3 clone RP11-79K12, WORKING DRAFT SEQUENCE,
14 unordered pieces.
ACCESSION AC076969
VERSION AC076969.6 GI:16117967
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 128118)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Aisbrook,S.L., Amaralung,H.C., Are,J.R., Banks,T., Barbata,J.,
Benton,U., Bimege,K., Blankenburg,K., Bonnin,D., Bouck,J.,
Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,
Burck,P., Burckett,C., Burrell,K.L., Byrd,N.C., Caron,T.F.,
Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,
Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C.,
Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,
Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,
Dean,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H.,
Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,
Elhaj,C., Escotto,M., Falls,T., Ferraruto,D., Flagg,N., Ford,J.,
Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,
Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S.,
Hamilton,K., Harris,C., Harris,K., Hart,M., Haylak,P., Hawes,A.,
Hernandez,J., Hernandez,O., Hodgson,A., Hogue,M., Hollaway,C.,
Hollins,B., Homsil,F., Howard,S., Huber,J., Huylk,S., Hume,J.,
Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolliver,S.,
Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J.,
Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,
Lewis,L., Li,U., Li,Z., Lichtarge,O., Lieu,C., Liu,C., Liu,W.,
Loussegod,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,
Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martindale,A.,
Martinez,E., Massey,E., Mawhinney,E., Mcleod,M.P., Meador,M.,
Mei,G., Metzker,M., Miner,G., Miner,D., Mitchell,T., Mohabbat,K.,
Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,N.,
Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokwenwo,S.,
Ogun,M., Okwou,G., Orazungu,N., Oviedo,R., Pace,A., Payton,B.,
Peery,J., Peter,L., Peters,L., Pickens,R., Prims,E., Pu,L.L.,
Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolle,M.,
Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shoostari,N.,
Stison,I., Sodergren,E., Sonalke,T., Sparks,A., Stanley,J.,
Stang,H., Sutton,A., Swalek,A., Tabor,P., Tamerisa,A., Tamerisa,K.,
Tang,H., Tansey,J., Taylor,C., Taylor,T., Teifod,B., Thomas,N.,
Thomas,S., Umanli,K., Vasquez,L., Vera,V., Villalob,D., Vinson,R.,
Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
Watlington,S., Williams,G., Williamson,A., Wleczyk,R., Wooden,S.,

```

```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 128118)
Submitted (01-NUC-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Oct 14, 2001 this sequence version replaced gi:10047573.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HBRJ
Center clone name: RP11-79K12
----- Summary Statistics
Sequencing vector: M13: L08821
Chemistry: Dye-terminator Big Dye: 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 111307 bases at least Q40
Consensus quality: 117905 bases at least Q30
Consensus quality: 121494 bases at least Q20
Estimated insert size: 122854; sum-of-coverage estimation
Quality coverage: 0x in Q20 bases; agarose-tp estimation
Quality coverage: 4x in Q20 bases; sum-of-coverage estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank/draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 14 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1
20354: contig of 20354 bp in length
20355 20454: gap of unknown length
20455 33071: contig of 12617 bp in length
33071 33171: gap of unknown length
33171 46935: contig of 13764 bp in length
46935 47035: gap of unknown length
47035 57066: contig of 10031 bp in length
57066 57166: gap of unknown length
57166 70378: contig of 13212 bp in length
70378 81786: gap of unknown length
81786 81887: contig of 11308 bp in length
81887 91429: gap of unknown length
91429 91528: contig of 9542 bp in length
91528 100453: gap of unknown length
100453 100555: contig of 8927 bp in length
100555 107879: gap of unknown length
107879 107979: contig of 7324 bp in length
107979 114149: gap of unknown length
114149 114245: contig of 6170 bp in length
114245 120186: gap of unknown length
120186 120286: contig of 5937 bp in length
120286 123236: gap of unknown length
123236 123336: contig of 2950 bp in length
123336 125606: gap of unknown length
125606 125706: contig of 2270 bp in length
125706 128118: gap of unknown length
128118 128118: contig of 2412 bp in length.
Location/Qualifiers
1. 128118
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP11-79K12"

```

FEATURES
source

1042	1065	743	758	1204	1224
2783	2808	3045	2963	2722	2723
7074	7158	1288	1262	321	<800
4087	3861	967	973	1066	1058
7976	7926	461	<800	470	<800
1233	1199	4649	4618	3055	3072
3875	3861	5289	5417	10961	10848
27	<800	1098	1095	6725	6888
1137	1065	798	809	1896	1903
750	749	111	<800	8682	8666
214	<800	7644	7609	4438	4342
836	887	10084	9932	514	<800
1670	1651	3641	3589	3077	3072
2650	2609	7555	7609	1371	1337
1890	1872			4185	4087
4615	4522			719	<800
1321	1273			4130	4087
				367	<800
				2979	3072
				333	<800

FEATURES

source

Location/Qualifiers
 1. 128583
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="3"
 /clone="RP11-79K17"
 /clone.lib="RP11 human BAC library 11"
 COUNT 37777 a 24752 c 24789 g 41265 t
 IN

Query Match 10.1%; Score 40; DB 9; Length 128583;
 Best Local Similarity 100.0%; Pred. No. 2.1e-12;
 Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 69 AAAACCTGGGCTCTGCTGCTGCTGCTGCTGCTCTCT 108
 DB 27283 AAAACCTGGGCTCTGCTGCTGCTGCTGCTGCTCTCT 27244

RESULT 6
 AC107979/c 148290 bp DNA linear PRI 01-JUL-2002
 LOCUS Homo sapiens chromosome 15, clone CTD-3049M7, complete sequence.
 DEFINITION
 AC107979
 VERSION AC107979.7 GI:21592043
 KEYWORDS
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 148290)
 Birren,B., Nusbaum,C. and Lander,E.

TITLE
JOURNAL
REFERENCE
AUTHORS

Homo sapiens chromosome 15, clone CTD-3049M7
 Unpublished
 2 (bases 1 to 148290)
 Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N.,
 Anderson,S., Baran,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
 Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
 Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
 Cooke,P., Dearrellano,K., Dewar,K., Diaz,J.S., Dodge,S., Fero,S.,
 Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
 Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
 Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
 Kamat,A., Karatas,A., Kells,C., Larocque,K., Lamazares,R.,
 Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C.,
 Macdonald,P., Major,J., Margulis,N., Matthews,C., McCarthy,M.,
 McKean,P., McKernan,K., Melidrim,J., Menus,L., Mhova,T.,
 Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C.,
 Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,
 Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
 Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
 Rosetti,M., Roy,A., Santos,R., Schauer,S., Schnupack,R., Seaman,S.,
 Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
 Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
 Topham,K., Travers,M., Travis,N., Triggillo,J., Vassiliev,H.,
 Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
 Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Submitted (24-JAN-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 148290)
 Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N.,
 Anderson,S., Baran,N., Bastien,V., Bloom,T., Boguslavsky,L.,
 Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
 Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
 Cook,A., Cooke,P., Dearrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
 Fero,S., Ferreira,P., Fitzgerald,M., Fitzhugh,W., Gage,D.,
 Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L.,
 Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,
 Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Larocque,K.,
 Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K.,
 Liu,G., Maclean,C., Macdonald,P., Major,J., Margulis,N.,
 Matthews,C., McCarthy,M., McKean,P., McKernan,K., Melidrim,J.,
 Menus,L., Mhova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
 Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,
 O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
 Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,
 Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S.,
 Schnupack,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
 Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S.,
 Theodore,J., Topham,K., Travers,M., Travis,N., Triggillo,J.,
 Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
 Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE
JOURNAL
REFERENCE
AUTHORS

Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 4 (bases 1 to 148290)
 Birren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S.,
 Baran,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
 Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
 Cook,A., Cooke,P., Dearrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
 Fero,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
 Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
 Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
 Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
 Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
 McCarthy,M., Melidrim,J., Menus,L., Mhova,T., Mlenga,V.,
 Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
 O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
 Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
 Roman,J., Roy,A., Schauer,S., Schnupack,R., Seaman,S., Severy,P.,
 Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
 Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
 Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
 Zembek,L., Zimmer,A. and Zody,M.

TITLE Direct Submission
JOURNAL Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT On Jun 26, 2002 this sequence version replaced gi:21321840.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L24533
Center clone name: 3049_M_7

FEATURES
Source Location/Qualifiers
1. 148290
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="CTD-3049M7"
/clone_lib="CTD2 Human BAC"
complement(1982..2092)
repeat_region
/rpt_family="MIR"
complement(3356..3654)
repeat_region
/rpt_family="AluSx"
3789..3844
/rpt_family="MER58"
complement(4429..4728)
repeat_region
/rpt_family="AluSp"
5494..5723
repeat_region
/rpt_family="(TA)n"
5727..5766
repeat_region
/rpt_family="(GA)n"
5806..5857
repeat_region
/rpt_family="AT-rich"
6377..6581
repeat_region
/rpt_family="L1"
6647..7069
repeat_region
/rpt_family="L1ME3"
7057..7631
repeat_region
/rpt_family="L1ME3"
complement(8821..8958)
repeat_region
/rpt_family="MIR3"
11129..11224
repeat_region
/rpt_family="FRAM"
11129..11338
repeat_region
/rpt_family="CT-rich"
11943..12090
repeat_region
/rpt_family="MIR"
12818..13060
repeat_region
/rpt_family="MIR"
13118..19146
repeat_region
/rpt_family="L1PA2"
20846..20912
repeat_region
/rpt_family="CT-rich"
21294..21314
repeat_region
/rpt_family="AT-rich"
21635..23417
repeat_region
/rpt_family="L1PA16"
23438..25940
repeat_region
/rpt_family="L1PA16"
26121..26145
repeat_region
/rpt_family="(TTCA)n"
26293..26417
repeat_region
/rpt_family="MIR"
complement(26872..26937)
repeat_region
/rpt_family="L2"
complement(27064..27431)
/rpt_family="L2"
27917..28357
repeat_region

repeat_region /rpt_family="MLT1B"
28460..28806
repeat_region /rpt_family="MLT1I"
29509..29698
repeat_region /rpt_family="MIR"
29716..29790
repeat_region /rpt_family="CT-rich"
29791..29811
repeat_region /rpt_family="AT-rich"
complement(31182..31787)
repeat_region /rpt_family="L2"
complement(32398..32581)
repeat_region /rpt_family="AluSg"
complement(32582..32777)
repeat_region /rpt_family="L1PB1"
complement(32778..32912)
repeat_region /rpt_family="AluSg"
33104..33207
repeat_region /rpt_family="L1MC5"
complement(33208..33365)
repeat_region /rpt_family="MER53"
33366..33529
repeat_region /rpt_family="L1MC5"
33877..34506
repeat_region /rpt_family="L1MED"
34667..34808
repeat_region /rpt_family="L1MED"
35585..36698
repeat_region /rpt_family="L1M4"
complement(36699..36830)
repeat_region /rpt_family="FLAM-C"
36831..38141
repeat_region /rpt_family="L1M4"
38194..38373
repeat_region /rpt_family="MLT1I"
39530..39757
repeat_region /rpt_family="MIR"
39992..40017
repeat_region /rpt_family="(TTTG)n"
40283..40583
repeat_region /rpt_family="AluY"
41414..41616
repeat_region /rpt_family="MLT1C"

Query Match 10.1%; Score 40; DB 9; Length 148290;
Best Local Similarity 100.0%; Pred. No. 2.1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 68 TAAACTCTGGGCTCTGTGTGCTGAGTGCGTCTC 107
Db 84499 TAAACATCCGCGGTCTGTGTGCTGAGTGCGTCTC 84460

RESULT 7
AC103996/c 165649 bp DNA linear PRI 01-JUL-2002
LOCUS
DEFINITION Homo sapiens chromosome 15, clone RP11-76E17, complete sequence.
ACCESSION AC103996
VERSION AC103996.7 GI:21637504
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE
1 (bases 1 to 165649)
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N.,
JOURNAL 2 (bases 1 to 165649)
REFERENCE Unpublished
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N.,
Anderson, S., Barina, N., Bastien, V., Boguslavsky, D., Boukhgalter, B.,
Brown, A., Camarata, J., Campio, A., Chang, J., Chazaro, B.,
Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,


```
repeat_region 19696..19750
/rpt_family="GA-rich"
repeat_region 19752..19920
/rpt_family="L1MD3"
repeat_region 19994..20102
/rpt_family="L2"
unsure 20049..20114
/note="single clone coverage"
repeat_region 20485..20655
/rpt_family="MIR3"
repeat_region complement(21285..21441)
/rpt_family="MIR"
repeat_region 21496..21717
/rpt_family="L2"
repeat_region complement(21720..21796)
/rpt_family="MIR"
repeat_region complement(21943..22131)
/rpt_family="MIR"
repeat_region 23082..23195
/rpt_family="L2"
repeat_region 23198..23248
/rpt_family="GA-rich"
repeat_region 23267..23333
/rpt_family="CATR)n"
repeat_region complement(24465..24833)
/rpt_family="MT1A2"
repeat_region 26142..26334
/rpt_family="MIR"
repeat_region complement(27659..27811)
/rpt_family="MIR"
repeat_region complement(28811..28880)
/rpt_family="MIR"
repeat_region complement(29026..29269)
/rpt_family="MIR"
repeat_region complement(29673..29706)
/rpt_family="MSTB"

Query Match 10.1%, Score 40; DB 9; Length 165649;
Best Local Similarity 100.0%; Pred. No. 2.1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 68 TAAACTCCTGGCTCTGTGTGTCCTGAGTGGCTGCTC 107
|||||
Db 8110 TAAACTCCTGGCTCTGTGTGTCCTGAGTGGCTGCTC 8071

RESULT 8
AC104303 178650 bp DNA linear PRI 25-FEB-2002
AC104303
AC104303 AC064830
AC104303.2 GI:18874945
HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 178650)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimmachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Unpublished
AUTHORS 2 (bases 1 to 178650)
Kaul,R.K., Olson,M.V., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Submitted (07-DEC-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (bases 1 to 178650)
Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimmachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Submitted (25-FEB-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
```

COMMENT

On Feb 25, 2002 this sequence version replaced gi:17402782.

----- Genome Center

Center: University of Washington Genome Center

Center Code: UWGC

Web site: <http://www.genome.washington.edu>

Contact: uwgchgs@u.washington.edu

Drafting Center: WUGSC

----- Project Information

Center project name: chr-3

Genet clone name: RP11-391P4 (bc0402)

----- Summary statistics

Sequencing vector: unknown; 55% of reads

Sequencing vector: plasmid; 45% of reads

Chemistry: Dye-terminator ET; 89% of reads

Chemistry: Dye-terminator Big Dye; 11% of reads

Assembly program: Phrap; version 0.990319

Consensus quality: 178494 bases at least Q40

Consensus quality: 178650 bases at least Q20

Insert size: 178648; sum-of-contigs

Quality coverage: 8.0x in Q20 bases; sum-of-contigs

Overlapping Sequences:

5': RP11-475023 (UWGC:bc0439) AC023346

3': RP11-79K12 AC076969

Sequence Quality Assessment:

This entry has been annotated with sequence quality

estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

Basebank flat file format but are available as part

of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:

all regions were either double-stranded or sequenced with an

alternate chemistry or covered by high quality data (i.e., Phred

quality >= 30); an attempt was made to resolve all sequencing

problems, such as compressions and repeats; all regions were

covered by at least one plasmid subclone or more than one M13

subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest

fingerprinting. Comparison of the experimentally derived digest

fragments with sequence-predicted fragments is given below.

The electronically-digested sequence consists of both insert and

vector, in order to accurately represent the entire circular BAC.

Small fragments below a variable cutoff (approximately 400-800 bp)

are not resolved in the fingerprint and hence do not appear

in the table. There are no significant remaining discrepancies

between the experimental and predicted values. Uniquely ordered

fragments are separated by dashed lines.

BgIII EcoRI HindIII

SeqDerMap FngPrnt SeqDerMap FngPrnt SeqDerMap FngPrnt

7050 7417 8696 8715 7163 7098

2067 2138 6 800 6382 6518

13472 13264 3077 3083 512 800

7644 7859 514 800 449 800

111 800 4438 4377 7988 7884

798 783 8682 8715 11779 11698

1098	1109	1896	1876	472	<800
5288	5395	6724	6948	783	787
4649	4507	10961	10773	926	926
461	<800	3055	3083	2431	2538
967	982	470	<800	1015	1029
1288	1268	1066	1037	758	<800
3045	2923	321	<800	1550	1541
743	783	2722	2683	361	<800
5628	5682	1204	1300	763	<800
2714	2699	6992	7281	2975	3002
4002	3900	7210	7830	3299	3468
227	<800	7821	8101	1907	1894
1794	1784	6855	6948	4020	4220
2168	2138	859	857	5158	5126
10032	9744	2629	2683	497	<800
6694	6810	1065	1037	4321	4220
1505	1491	3934	3965	887	926
3312	3304	859	857	180	<800
896	912	931	935	4699	4887
5873	5938	858	857	4755	4674
78	<800	799	857	26	<800
2135	2138	1895	1876	906	926
133	<800	58	<800	2325	2346
11939	11575	5129	5076	369	<800
1507	1491	6051	6048	2523	2538
8069	8181	3538	3537	1889	1894
1632	1617	1114	1094	4437	4428
3729	3678	8420	8391	373	<800
5533	5395	1024	1037	24	<800
358	<800	1323	1380	263	<800
3764	3678	1502	1481	632	<800
7022	6810	4538	4539	1258	1247
8534	8821	1414	1380	1114	1083
5206	5050	521	<800	1370	1320
9052	9744	8346	8391	1262	1247
1625	1617	39	<800	1546	1541
749	783	2716	2683	459	<800

Query Match 10.1%; Score 40; DB 9; Length 178650;
 Best Local Similarity 100.0%; Pred. No. 2,1e-12;
 Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 69 AAACTCCTGGCTCTGTGTGCTGAGTGGCTGCTCT 108
 Db 165449 AAACTCCTGGCTCTGTGTGCTGAGTGGCTGCTCT 165488

RESULT 9
 AC090762/c 192826 bp DNA linear PRI 28-FEB-2002
 LOCUS Homo sapiens chromosome 15, clone RP11-387E8, complete sequence.
 DEFINITION AC090762
 AC090762.9 GI:18997378
 VERSION
 KEYWORDS
 SOURCE Homo sapiens.
 ORGANISM Homo sapiens.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
 1 (bases 1 to 192826)
 Birren,B., Linton,L., Nusbaum,C. and Lander,E.
 Unpublished
 2 (bases 1 to 192826)
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
 Barre,N., Bastien,V., Boguslavsky,L., Boukhalter,B., Brown,A.,
 Camarata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,
 Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
 Dodge,S., Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,
 Gardina,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,
 Hagos,B., Heaford,A., Horton,L., Hulme,M., Iliev,I., Johnson,R.,
 Jones,C., Karafas,A., Larocque,K., Lamazares,R., Landers,T.,
 Lehoccky,J., Levine,R., Liu,G., Maclean,C., Macdonald,P.,
 Marquis,N., Mathews,C., McCarthy,M., McEwan,P., McKernan,K.,
 McPheters,R., Meldrim,J., Menus,L., Mihova,T., Mienga,V.,
 Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
 O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
 Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R.,
 Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M.,
 Roy,A., Santos,R., Schauer,S., Schnupack,R., Seaman,S., Severy,P.,
 Sougnaz,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
 Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,

Trevors, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
Submitted Submission

JOURNAL
Submitted (10-MAR-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

REFERENCE
3 (bases 1 to 192826)

AUTHORS
Birren, B., Linton, L., Nusbaum, C., Lander, E., All, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhgalter, B., Brown, A., Camarata, J., Campiolo, A., Chang, J., Chazaro, B., Choquel, Y., Collangelo, M., Collins, S., Collamore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N., Hago, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Larocque, K., Lamazares, R., Landers, T., Lehocsky, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Melidim, J., Meneses, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, D., Testaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
Submitted Submission

JOURNAL
Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT
On Feb 28, 2002 this sequence version replaced g1:18377189. All repeats were identified using RepeatMasker:

Smith, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

TITLE
Genome Center

JOURNAL
Center: Whitehead Institute/ MIT Center for Genome Research

COMMENT
Center code: WIBR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu

----- Project Information -----
Center project name: L12392
Center clone name: 387_E_8

----- Location/Qualifiers -----
1. 192826
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-387E8"
/clone_lib="RP11 Human Male BAC"
complement(864..1222)
/rpt_family="L1PB3"
1223..1252
/rpt_family="(TAGG)n"
complement(1253..1492)
/rpt_family="L1PB3"
4287..4576
/rpt_family="AluDb"
4671..5034
/rpt_family="THE1B"
5130..5438
/rpt_family="AluSx"
5517..5662
/rpt_family="MIR"
5798..3827
/rpt_family="AT_rich"
5985..6114
/rpt_family="AluDb"
6227..6253
/rpt_family="(CA)n"

repeat_region 7329..7446
/rpt_family="MIR"
complement(8147..8452)
repeat_region
/rpt_family="AluDb"
8540..8639
/rpt_family="MER45"
complement(8832..8916)
repeat_region
/rpt_family="L2"
10391..10599
/rpt_family="MER3"
11688..11828
/rpt_family="L1MC/D"
12029..12078
/rpt_family="AT_rich"
12092..12447
/rpt_family="THE1C"
complement(13616..13751)
/rpt_family="MIR3"
complement(13958..14142)
/rpt_family="MIR"
complement(15277..15553)
/rpt_family="MER8"
complement(15811..16005)
/rpt_family="MIR"
complement(16101..16440)
/rpt_family="L3"
16920..16958
/rpt_family="(TCCC)n"
complement(17145..17444)
/rpt_family="AluSx"
complement(18418..19953)
/rpt_family="L1MEC"
complement(19978..20262)
/rpt_family="L1MEC"
complement(20288..20794)
/rpt_family="L1MEC"
complement(20822..21097)
/rpt_family="L1MEC"
complement(21345..21743)
/rpt_family="L1MEC"
22599..23518
/rpt_family="L1MEC"
23527..23901
/rpt_family="L1MEC"
complement(23927..24026)
/rpt_family="MSTB1"
complement(24027..24256)
/rpt_family="MER30"
complement(24257..24557)
/rpt_family="MSTB1"
24563..24594
/rpt_family="AT_rich"
complement(25258..25639)
/rpt_family="L1MC"
complement(25688..25838)
/rpt_family="L1MC"
26167..26506
/rpt_family="THE1B"
26746..27094
/rpt_family="Tiger2a"
27095..27184
/rpt_family="MADE1"
27185..27279
/rpt_family="Tiger2a"
28321..28363
/rpt_family="(TG)n"
29172..29333
/rpt_family="MIR"
30664..30898
/rpt_family="L1MB8"
31577..31598
/rpt_family="AT_rich"
32378..32475
repeat_region

Query Match 10.1% Score 40; DB 9; Length 192826;
 Best Local Similarity 100.0%; Pred. No. 2;le-12;
 Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0.

Oy 68 TAAACCTCCTGGCTCTGTGTGTGCTAGTGGCTGCTC 107
 |||||||
 Db 190946 TAAACCTCCTGGCTCTGTGTGTGCTAGTGGCTGCTC 190907

RESULT 10
 LOCUS AC007445 32918 bp DNA linear HTG 30-JUN-2000
 DEFINITION Homo sapiens chromosome 18 clone RP11-344B7 map 18, *** SEQUENCING
 AC007445
 ACCESSION AC007445
 VERSION AC007445.5 GI:8844149
 KEYWORDS HTG; HTGS_PHASE2.
 SOURCE Homo sapiens.
 ORGANISM Homo sapiens.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
 1 (bases 1 to 32918)
 Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 Homo sapiens chromosome 18, clone RP11-344B7
 Unpublished
 2 (bases 1 to 32918)
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, M.,
 Baker, J., Baldwin, J., Barna, N., Beckert, R., Benn, J., Brown, A.,
 Castle, A., Cerny, J., Colangelo, M., Collins, S., Collumore, A.,
 Cooke, P., DeRubeis, K., Depierre, E., Devon, K., Dewar, K.,
 Donelan, L., Doyle, M., Ferreira, P., Fitzhugh, W., Forrest, C.,
 Funke, R., Gage, D., Galagan, J., Gardyna, S., Gilbert, D., Grant, G.,
 Hagos, B., Heaford, A., Horton, L., Howland, J. C., Jones, C., Kuan, L.,
 Karats, A., Lechoczky, J., Lien, C., Locke, K., MacDonald, P.,
 Marquis, N., McEwan, P., McGurk, A., McKernan, K., McLaughlin, J.,
 Meldrum, J., Molla, M., Morris, W., Morrow, J., Mychaleckyj, J.,
 Naylor, J., Niloff, M., O'Connor, T., O'Donnell, P., Pavlin, B.,
 Peterson, K., Pollara, V., Riley, R., Roberts, D., Roy, A., Severy, P.,
 Stange-Thomann, N., Stojanovic, N., Stone, C., Subramanian, A.,
 Tefaye, S., Toruella-Miller, I., Vassiliev, H., Vo, A., Wagner, A.,
 Wheeler, J., Wu, X., Wyman, D., Ye, W. J. and Zody, M.
 Direct Submission

TITLE
 JOURNAL
 COMMENT
 Submitted (30-APR-1999) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Jun 30, 2000 this sequence version replaced gi:8705092.
 All repeats were identified using RepeatMasker:
 Smit, A. F. A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: MIBR	
Web site: http://www.seq.wi.mit.edu	
Contact: sequence_submissions@genome.wi.mit.edu	
----- Project Information	
Center project name: L571	
Center clone name: 344_B_7	

* NOTE: This is a 'working draft' sequence. It currently	
* consists of 1 contigs. Gaps between the contigs	
* are represented as runs of N. The order of the pieces	
* is believed to be correct as given, however the sizes	
* of the gaps between them are based on estimates that have	
* provided by the submittor.	
* This sequence will be replaced	
* by the finished sequence as soon as it is available and	
* the accession number will be preserved.	
* 1 32918: contig of 32918 bp in length.	
Location/Qualifiers	
1. 32918	
/organism="Homo sapiens"	
/db_xref="taxon:9606"	
/chromosome="18"	
/map="18"	
/clone="RP11-344B7"	
/clone_1ib="RPC1-11 Human Male BAC"	
9462 a 6493 c 6865 g 9853 t 245 others	
BASE COUNT	
ORIGIN	
Query Match 9.8%; Score 39; DB 2; Length 32918;	
Best Local Similarity 100.0%; Prid. No. 8.2e-12;	
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Oy 60 GGAGTATGTAACACCTCGGCTCTGTGTGTCGCTGAG 98	
Db 3535 GGAGTATGTAACACCTCGGCTCTGTGTGTCGCTGAG 3573	
RESULT 11	
AL358817	
LOCUS	
DEFINITION	
Human DNA sequence from clone Rp11-399N22 on chromosome 10,	
complete sequence.	
AL358817	
AL358817.18 GI:15990637	
HTG.	
SOURCE	
ORGANISM	
human.	
Homo sapiens	
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
1 (bases 1 to 38936)	
Lovell,J.	
Direct Submission	
Submitted (06-OCT-2001) Sanger Centre, Hinxton, Cambridgeshire,	
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk	
requests: clonerequests@sanger.ac.uk	
On Oct 9, 2001 this sequence version replaced gi.14669268.	
During sequence assembly data is compared from overlapping clones.	
Where differences are found these are annotated as variations	
together with a note of the overlapping clone name. Note that the	
variation annotation may not be found in the sequence submission	
corresponding to the overlapping clone, as we submit sequences with	
only a small overlap as described above.	
This sequence was finished as follows unless otherwise noted: all	
regions were either double-stranded or sequenced with an alternate	
chemistry or covered by high quality data (i.e., phred quality >=	
30); an attempt was made to resolve all sequencing problems, such	
as compressions and repeats; all regions were covered by at least	
one plasmid subclone or more than one M13 subclone; and the	
assembly was confirmed by restriction digest. The following	
abbreviations are used to associate primary accession numbers given	
in the feature table with their source databases: Em., EMBL; Sw.,	
SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP	
database can be found at	

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 10, constructed by the Sanger Centre Chromosome 10 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/chr10>

RP11-399N22 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/dacpac/home.htm>

VECTOR: PBAC3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-399N22. It may be shorter because we sequence overlapping sections only once, except for a short overlap. The true left end of clone RP11-432J9 is at 36937 in this sequence. The true right end of clone RP11-91A1 is at 2000 in this sequence.

FEATURES

source

1.38936
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-399N22"
/clone_1lb="RP11-11.2"
9315 a 9079 c 9111 g 11431 t
ORIGIN

Query Match 9.8%; Score 39; DB 9; Length 38936;
Best Local Similarity 100.0%; Pred. No. 8.2e-12;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Caps 0;

QY 60 GGAGTATGTAAACCTCGGTCTCTGTCGTGCTGAG 98
|||||
Db 28896 GGAGTATGTAAACCTCGGTCTCTGTCGTGCTGAG 28934

RESULT 12

AC025179

LOCUS AC025179 124271 bp DNA linear HTG 20-APR-2001
DEFINITION Homo sapiens chromosome 5 clone CTD-2174B5, WORKING DRAFT SEQUENCE,
8 unordered pieces.

AC025179

AC025179.4 GI:13699647

HTG: HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEPIN.

KEYWORDS

SOURCE

ORGANISM

Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1052: contig of 1052 bp in length
* 1053 1152: gap of unknown length
* 1153 3331: contig of 2179 bp in length
* 3332 3431: gap of unknown length
* 3432 7370: contig of 3939 bp in length
* 7371 7470: gap of unknown length
* 7471 15483: contig of 8013 bp in length
* 15484 15583: gap of unknown length
* 15584 24916: contig of 9333 bp in length
* 24917 25016: gap of unknown length
* 25017 39922: contig of 14506 bp in length
* 39923 40022: gap of unknown length
* 40023 68684: contig of 28662 bp in length
* 68685 68785: gap of unknown length
* 68785 124271: contig of 55487 bp in length.

FEATURES

source

1.124271
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2174B5"
/clone_1lb="Caltech human BMC library D"
36863 a 24233 c 23599 g 38876 t 700 others
ORIGIN

Query Match 9.8%; Score 39; DB 2; Length 124271;
Best Local Similarity 100.0%; Pred. No. 8.6e-12;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Caps 0;

QY 61 GAGTATGTAAACCTCGGTCTCTGTCGTGCTGAGT 99
|||||
Db 79646 GAGTATGTAAACCTCGGTCTCTGTCGTGCTGAGT 79684

RESULT 13

AC008814

LOCUS AC008814 146671 bp DNA linear PRI 31-OCT-2001
DEFINITION Homo sapiens chromosome 5 clone CTD-2117L12, complete sequence.

AC008814

AC008814.6 GI:16554342

HTG:

KEYWORDS

SOURCE

ORGANISM

Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

JOURNAL

Project Information
Center Project Name: 694394
Center clone name: CTRB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently

Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Oct 31, 2001 this sequence version replaced gi:15290309.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing completed at Stanford Human Genome Center
www.shgc.stanford.edu
Quality: Phrap Quality >=40 99.3% of Sequence;
Estimated Total Number of Errors is 0.7.

FEATURES
Source Location/Qualifiers
1.146671
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2117J12"

BASE COUNT 44951 a 28815 c 28434 g 44471 t
ORIGIN

Query Match 9.8% Score 39; DB 9; Length 146671;
Best Local Similarity 100.0%; Pred. No. 8.6e-12;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 61 GAGTATGTAACCTCGTGGTCTCTGTGTGTCCTGAGT 99
|||||
Db 54480 GAGTATGTAACCTCGTGGTCTCTGTGTGTCCTGAGT 54518
|||||

RESULT 14
AP001019/c
AP001019 159747 bp DNA linear HTG 30-MAY-2000
Homo sapiens chromosome 18 clone RP11-752111 map 18p11.3, WORKING
DRAFT SEQUENCE, 24 unordered pieces.

ACCESSION AP001019.2 GI:8117689
VERSION AP001019
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens DNA, clone:RP11-752111.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 159747)
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE Homo sapiens 159,747 genomic DNA of 18p11.3
JOURNAL Published only in Database (2000)
REFERENCE 2 (bases 1 to 159747)
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE Direct Submission
JOURNAL Submitted (05-JAN-2000) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gsc.riken.go.jp,
URL:http://hnp.gsc.riken.go.jp/, Tel:81-42-778-9923,
Fax:81-42-778-9924)
COMMENT On May 31, 2000 this sequence version replaced gi:6997769.
----- Genome Center
Center: RIKEN Genomic Sciences Center(GSC)
Center code: RIKEN
Web site: http://hnp.gsc.riken.go.jp/
Contact: hattori@gsc.riken.go.jp
----- Project Information
Center project name: Humdraft18
Center clone name: RP11-752111
----- Summary Statistics
Sequencing vector: PCR products; 100% of reads
Chemistry: Dye-terminator; EF-amersham; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 145356 bases at least Q40
Consensus quality: 152227 bases at least Q30
Consensus quality: 155843 bases at least Q20
Insert size: 157447; sum-of-ctrls
Quality coverage: 4.51x in Q20 bases; sum-of-ctrls

NOTE: This is a 'working draft' sequence. It currently consists of
24 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved
1 18444 contig of 18444 bp in length
18545 29631 contig of 11087 bp in length

29732 43143 contig of 13412 bp in length
43244 57170 contig of 13927 bp in length
57271 68240 contig of 10970 bp in length
68341 78134 contig of 9794 bp in length
78235 87873 contig of 9639 bp in length
87974 96197 contig of 8224 bp in length
96298 103976 contig of 6650 bp in length
104077 110726 contig of 6500 bp in length
110827 116866 contig of 6040 bp in length
116867 116966 contig of 100 bp in length
116967 122789 contig of 5823 bp in length
122790 122889 contig of 100 bp in length
122890 127665 contig of 4776 bp in length
127666 127765 contig of 100 bp in length
127766 132618 contig of 4853 bp in length
132619 132718 contig of 100 bp in length
132719 136051 contig of 3333 bp in length
136052 136151 contig of 100 bp in length
136152 139749 contig of 3598 bp in length
139750 139849 contig of 100 bp in length
139850 142810 contig of 2961 bp in length
142811 142910 contig of 100 bp in length
142911 143855 contig of 2945 bp in length
143856 145955 contig of 100 bp in length
145956 148840 contig of 2885 bp in length
148841 148940 contig of 100 bp in length
148941 151243 contig of 2303 bp in length
151244 151347 contig of 100 bp in length
151348 153454 contig of 2111 bp in length
153455 153554 contig of 100 bp in length

Sequence updated (26-May-2000).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 24 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 18444: contig of 18444 bp in length
18445 18544: gap of 100 bp
18545 29631: contig of 11087 bp in length
29632 29731: gap of 100 bp
29732 43143: contig of 13412 bp in length
43144 43243: gap of 100 bp
43244 57170: contig of 13927 bp in length
57171 57270: gap of 100 bp
57271 68240: contig of 10970 bp in length
68241 68340: gap of 100 bp
68341 78134: contig of 9794 bp in length
78135 78234: gap of 100 bp
78235 87873: contig of 9639 bp in length
87874 87973: gap of 100 bp
87974 96197: contig of 8224 bp in length
96198 96297: gap of 100 bp
96298 103976: contig of 7679 bp in length
103977 104076: gap of 100 bp
104077 110726: contig of 6650 bp in length
110727 110826: gap of 100 bp
110827 116866: contig of 6040 bp in length
116867 116966: gap of 100 bp
116967 122789: contig of 5823 bp in length
122790 122889: gap of 100 bp
122890 127665: contig of 4776 bp in length
127666 127765: gap of 100 bp
127766 132618: contig of 4853 bp in length
132619 132718: gap of 100 bp
132719 136051: contig of 3333 bp in length
136052 136151: gap of 100 bp
136152 139749: contig of 3598 bp in length
139750 139849: gap of 100 bp
139850 142810: contig of 2961 bp in length
142811 142910: gap of 100 bp
142911 143855: contig of 2945 bp in length
143856 145955: gap of 100 bp
145956 148840: contig of 2885 bp in length
148841 148940: gap of 100 bp
148941 151243: contig of 2303 bp in length
151244 151347: gap of 100 bp
151348 153454: contig of 2111 bp in length
153455 153554: gap of 100 bp

* 153555 156134: contig of 2580 bp in length
* 156135 156234: gap of 100 bp
* 156235 158089: contig of 1855 bp in length
* 158090 158189: gap of 100 bp
* 158190 159747: contig of 1558 bp in length.

FEATURES
Location/Qualifiers

source

1. 159747

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="18"

/map="18p11.3"

/clone="RP11-752111"

1. 18444

/note="assembly_fragment"

18545. .29631

/note="assembly_fragment"

29732. .43143

/note="assembly_fragment"

43244. .57170

/note="assembly_fragment clone_end:SP6 vector_side:left"

57271. .68240

/note="assembly_fragment"

68341. .78134

/note="assembly_fragment"

78235. .87873

/note="assembly_fragment"

87974. .96197

/note="assembly_fragment"

96298. .103976

/note="assembly_fragment clone_end:T7 vector_side:left"

104077. .110726

/note="assembly_fragment"

110827. .116866

/note="assembly_fragment"

116967. .122789

/note="assembly_fragment"

122890. .127665

/note="assembly_fragment"

127766. .132618

/note="assembly_fragment"

132719. .136051

/note="assembly_fragment"

136132. .139749

/note="assembly_fragment"

139850. .142810

/note="assembly_fragment"

142911. .145855

/note="assembly_fragment"

145956. .148840

/note="assembly_fragment"

148941. .151243

/note="assembly_fragment"

151344. .153454

/note="assembly_fragment"

153555. .156134

/note="assembly_fragment"

156235. .158089

/note="assembly_fragment"

158190. .159747

/note="assembly_fragment"

159747

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

162740

/note="assembly_fragment"

BASE COUNT 47302 a 30397 c 29897 g 49849 t 2302 others
ORIGIN

Query Match 9.8%: Score 39; DB 2; Length 159747;

Best Local Similarity 100.0%; Pred. No. 8 6e-12;

Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 60 GGAGTATGTAAACTCGGCTCTCTGTGTGCTGCTGAG 98

DB 74932 GGAGTATGTAAACTCGGCTCTCTGTGTGCTGCTGAG 74894

RESULT 15

AC034249

LOCUS

DEFINITION

AC034249

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

AC034249 162740 bp DNA linear HTG 31-AUG-2001

Homo sapiens chromosome 5 clone RP11-427C17, WORKING DRAFT

SEQUENCE, 6 ordered pieces.

AC034249

AC034249.3 GI:15383785

HTG: HTGS_PHASE2; HTGS_DRAFT; HTGS_ACTIVEFTN.

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 162740)

DOE Joint Genome Institute.

Sequencing of Human Chromosome 5

Unpublished

2 (bases 1 to 162740)

DOE Joint Genome Institute.

Direct Submission

Submitted (05-APR-2000) Production Sequencing Facility, DOE Joint

Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA

On Aug 31, 2001 this sequence version replaced gi:9211234.

-----Genome Center

Center: Joint Genome Institute

Center Code: JGI

Web site: <http://www.jgi.doe.gov>

Project Information

Center Project Name: 570398

Center clone name: RPCT-11_427C17

Summary Statistics

Consensus quality: 158733 bases at least Q40

Consensus quality: 161469 bases at least Q30

Consensus quality: 162087 bases at least Q20

Estimated insert size: 160000; pulse field gel estimation

Estimated insert size: 162240; sum-of-contigs estimation

Quality coverage: 9.28 in Q20 bases; pulse field gel estimation

Quality coverage: 9.16 in Q20 bases; sum-of-contigs estimation.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 6 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submitter.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

1 85871: contig of 85871 bp in length

* 85872 85971: gap of unknown length in length

* 85972 96765: contig of 10794 bp in length

* 96766 96865: gap of unknown length

* 96866 103302: contig of 6437 bp in length

* 103303 103402: gap of unknown length

* 103403 111971: contig of 8569 bp in length

* 111972 112071: gap of unknown length in length

* 112072 129034: contig of 16563 bp in length

* 129035 129134: gap of unknown length

* 129135 162740: contig of 33606 bp in length.

Location/Qualifiers

1. 162740

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="5"

/clone="RP11-427C17"

/clone.lib="RPCT human BAC library 11"

BASE COUNT 48722 a 31435 c 31563 g 50520 t 500 others

ORIGIN

Query Match 9.8%: Score 39; DB 2; Length 162740;

Best Local Similarity 100.0%; Pred. No. 8.7e-12;

Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 61 GAGTAGTAAACTCGGCTCTCTGTGTGCTGCTGAGT 99

Mon Apr 28 09:26:00 2003

us-09-513-999c-3792_copy_51_446.oli.rge

Page 16

Db 3599 GAGTATGTAAACCTCTGGGTCTCTGTGTGTGCTGAGT 3637

Search completed: April 25, 2003, 00:41:44
Job time : 1748.12 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:19:38 ; Search time 181.988 Seconds
(without alignments)
4900.271 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atgggtgatcttttgcctt.....gamctgatattcattga 396

Scoring table:

OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 2185239 segs, 112599159 residues

Word size : 0

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N.Geneseq_101002.*

1: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
2: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
3: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
4: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
5: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
6: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
7: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
8: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
9: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
10: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
11: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
12: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
13: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
14: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
15: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
16: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
17: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
18: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
19: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
20: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
21: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SIDs2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	396	100.0	447	21 AAC03794	Human secreted pro
2	396	100.0	447	21 AAZ42680	Human 5' EST isola
3	29	7.3	966	23 AAS93299	DNA encoding novel
4	24	6.1	660	23 AAS73441	DNA encoding novel
5	22	5.6	3659	7 AAN60204	Interferon-pseudo-
6	22	5.6	147724	24 ABR83566	Human cDNA differe
7	20	5.1	1179	23 AAS91061	DNA encoding novel
8	20	5.1	2526	23 AAS66674	DNA encoding novel
9	19	4.8	366	22 AAC90702	Human secretory pr

C	10	19	4.8	802	20 AAX84949
C	11	19	4.8	1982	21 AAC68089
C	12	19	4.8	2770	23 ABL09112
C	13	19	4.8	4902	23 ABL28930
C	14	18	4.5	31	16 AAT45064
C	15	18	4.5	339	21 AAZ43080
C	16	18	4.5	431	24 ABR24383
C	17	18	4.5	431	24 ABR24390
C	18	18	4.5	432	22 AAI83964
C	19	18	4.5	570	22 AAB63453
C	20	18	4.5	570	22 ABA30652
C	21	18	4.5	570	22 AAK11985
C	22	18	4.5	570	22 AAK37688
C	23	18	4.5	570	22 AAI18447
C	24	18	4.5	570	22 AAI43563
C	25	18	4.5	570	22 AAS11680
C	26	18	4.5	827	23 AAS87115
C	27	18	4.5	852	22 AAI94104
C	28	18	4.5	870	22 AAI94266
C	29	18	4.5	872	21 AAC79760
C	30	18	4.5	3342	23 AAS87118
C	31	18	4.5	4886	22 AAK85823
C	32	18	4.5	6464	22 ABA09665
C	33	18	4.5	9899	22 AAK85825
C	34	18	4.5	10918	22 ABA09581
C	35	18	4.5	10920	22 ABA09581
C	36	18	4.5	113515	24 ABL34174
C	37	17	4.3	266	20 AAV89709
C	38	17	4.3	288	21 AAC70647
C	39	17	4.3	288	21 AAC70650
C	40	17	4.3	288	21 AAC70656
C	41	17	4.3	288	21 AAC70668
C	42	17	4.3	288	21 AAC70671
C	43	17	4.3	294	22 AAH73108
C	44	17	4.3	297	22 AAH70408
C	45	17	4.3	318	22 AAH69668

ALIGNMENTS

RESULT 1	
AAC03794	standard; cDNA; 447 BP.
ID AAC03794	
AC AAC03794	
XX	
XX	06-OCT-2000 (first entry)
XX	
XX	Human secreted protein 5' EST, SEQ ID NO: 3792.
DE	
DE	Human; 5' EST: expressed sequence tag; secreted protein; cDNA isolation;
XX	gene therapy; chromosome mapping; ss.
KW	
KW	Human sapiens.
XX	
OS	Homo sapiens.
XX	
PN	EP103401-A2.
XX	
XX	06-SEP-2000.
PD	
XX	
XX	21-FEB-2000; 2000EP-0200610.
PF	
XX	26-FEB-1999; 99US-0122487.
PR	
XX	
PA	(GEST) GENSET.
PI	Dumas Milne Edwards J, Duclert A, Giordano J;
XX	
XX	WPI: 2000-500381/45.
DR	P-PSDB; AAG03788.
XX	
XX	New nucleic acid that is a 5' expressed sequence tag (5' EST) for
PT	obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for

RESULT 3
 AAS93299 ID AAS93299 standard; cDNA: 966 BP.
 XX AC AAS93299;
 XX DT 13-FEB-2002 (first entry)
 XX DE DNA encoding novel human diagnostic protein #29103.
 XX KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
 XX KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX OS Homo sapiens.
 XX PN WO200175067-A2.
 XX PD 11-OCT-2001.
 XX PF 30-MAR-2001; 2001WO-US08631.
 XX PR 31-MAR-2000; 2000US-0540217.
 XX PR 23-AUG-2000; 2000US-0649167.
 XX PA (HYSE-) HYSEQ INC.
 XX PI Drmanac RT, Liu C, Tang YT;
 XX DR WPI: 2001-639362/73.
 XX DR P-PSDB: ABG29112.
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS Claim 1; SEQ ID No 29103; 103pp; English.
 XX CC The invention relates to isolated polynucleotide (I) and
 XX CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 XX CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 XX CC and gene mapping, and in recombinant production of (II). The
 XX CC polynucleotides are also used in diagnostics as expressed sequence tags
 XX CC for identifying expressed genes. (I) is useful in gene therapy techniques
 XX CC to restore normal activity of (II) or to treat disease states involving
 XX CC (II). (II) is useful for generating antibodies against it, detecting or
 XX CC quantitating a polypeptide in tissue, as molecular weight markers and as
 XX CC a food supplement. (II) and its binding partners are useful in medical
 XX CC imaging of sites expressing (II). (I) and (II) are useful for treating
 XX CC disorders involving aberrant protein expression or biological activity.
 XX CC The polypeptide and polynucleotide sequences have applications in
 XX CC diagnostics, forensics, gene mapping, identification of mutations
 XX CC responsible for genetic disorders or other traits to assess biodiversity
 XX CC and to produce other types of data and products dependent on DNA and
 XX CC amino acid sequences. AAS64197-AAS94564 represent novel human
 XX CC diagnostic coding sequences of the invention.
 XX CC Note: The sequence data for this patent did not appear in the printed
 XX CC specification, but was obtained in electronic format directly from WIPO
 XX CC at ftp.wipo.int/pub/published_pct_sequences.
 XX SO Sequence 966 BP; 250 A; 221 C; 259 G; 236 T; 0 other;
 XX
 Query Match 7.3%; Score 29; DB 23; Length 966;
 Best Local Similarity 100.0%; Pred. No. 0.00027;
 Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 197 CAAAGATCTGTGGAGAAAGTGTGTTCC 225
 ||||||||||||||||||||||||||||
 Db 211 CAAAGATCTGTGGAGAAAGTGTGTTCC 239

RESULT 4
 AAS73441/C ID AAS73441 standard; cDNA: 660 BP.
 XX AC AAS73441;
 XX DT 13-FEB-2002 (first entry)
 XX DE DNA encoding novel human diagnostic protein #9245.
 XX KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
 XX KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX OS Homo sapiens.
 XX PN WO200175067-A2.
 XX PD 11-OCT-2001.
 XX PF 30-MAR-2001; 2001WO-US08631.
 XX PR 31-MAR-2000; 2000US-0540217.
 XX PR 23-AUG-2000; 2000US-0649167.
 XX PA (HYSE-) HYSEQ INC.
 XX PI Drmanac RT, Liu C, Tang YT;
 XX DR WPI: 2001-639362/73.
 XX DR P-PSDB: ABG09254.
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS Claim 1; SEQ ID No 9245; 103pp; English.
 XX CC The invention relates to isolated polynucleotide (I) and
 XX CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 XX CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 XX CC and gene mapping, and in recombinant production of (II). The
 XX CC polynucleotides are also used in diagnostics as expressed sequence tags
 XX CC for identifying expressed genes. (I) is useful in gene therapy techniques
 XX CC to restore normal activity of (II) or to treat disease states involving
 XX CC (II). (II) is useful for generating antibodies against it, detecting or
 XX CC quantitating a polypeptide in tissue, as molecular weight markers and as
 XX CC a food supplement. (II) and its binding partners are useful in medical
 XX CC imaging of sites expressing (II). (I) and (II) are useful for treating
 XX CC disorders involving aberrant protein expression or biological activity.
 XX CC The polypeptide and polynucleotide sequences have applications in
 XX CC diagnostics, forensics, gene mapping, identification of mutations
 XX CC responsible for genetic disorders or other traits to assess biodiversity
 XX CC and to produce other types of data and products dependent on DNA and
 XX CC amino acid sequences. AAS64197-AAS94564 represent novel human
 XX CC diagnostic coding sequences of the invention.
 XX CC Note: The sequence data for this patent did not appear in the printed
 XX CC specification, but was obtained in electronic format directly from WIPO
 XX CC at ftp.wipo.int/pub/published_pct_sequences.
 XX SO Sequence 660 BP; 149 A; 127 C; 148 G; 236 T; 0 other;
 XX
 Query Match 6.1%; Score 24; DB 23; Length 660;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 69 AAACCTCTGGGTCTGTGTGTG 92
 ||||||||||||||||||||||||
 Db 656 AAACCTCTGGGTCTGTGTGTG 633
 RESULT 5
 AAN60204

ID AAN60204 standard; DNA; 3659 BP.
 XX AAN60204;
 AC
 XX
 DT 05-AUG-1991 (first entry)
 XX
 DE Interferon-pseudo-omega-4.
 XX
 KW Interferon-omega; virucide; antitumor; SS.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT CDS 2951..3251
 FT /*tag= a
 FT /label= interferon-pseudo-omega-4
 XX
 PN EPI70204-A.
 XX
 XX 05-FEB-1986.
 XX
 XX 24-JUL-1985; 85EP-0109284.
 XX
 XX 14-FEB-1985; 85DE-3505060.
 XX 01-AUG-1984; 84DE-3428370.
 XX
 XX (BOEH) BOEHRINGER INGELHEIM.
 XX
 XX Hauptmann R, Meindl P, Dworkin-Rastl E, Adolf G, Sweetly P;
 PI Pieler C, Hanel N;
 XX
 XX WPI: 1986-036962/06.
 XX
 XX P-PSDB: AAP60256.
 XX
 XX New interferon omega polypeptide derivs. - useful as antiviral
 PT and antitumor agents, and new DNA sequences and genes coding for
 PT them.
 XX
 XX PS Disclosure; Fig 14; 115pp; German.
 XX
 XX The sequence encodes an interferon analogue, which has virucide
 CC and antitumor activity, and shows a synergistic increase in
 CC activity with interferon-gamma.
 CC
 XX Sequence 3659 BP; 796 A; 885 C; 1039 G; 939 T; 0 other;
 SQ
 Query Match 5.6%; Score 22; DB 7; Length 3659;
 Best Local Similarity 100.0%; Pred. No. 0.7;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 194 TTGCAAGATCTGTGAGAG 215
 ||||||||||||||||||
 Db 2557 TTGCAAGATCTGTGAGAG 2578
 RESULT 6
 ABR83566/c
 ID ABR83566 standard; cDNA; 147724 BP.
 XX
 AC ABR83566;
 XX
 DT 14-AUG-2002 (first entry)
 XX
 DE Human cDNA differentially expressed in granulocytic cells #137.
 XX
 KW Human; SS; granulocytic cell; DNA chip; bacterial infection;
 KW viral infection; parasitic infection; protozoal infection;
 KW fungal infection; sterile inflammatory disease; psoriasis;
 KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
 KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.

XX Homo sapiens.
 OS
 XX WC0200228999-A2.
 PN
 XX
 PD 11-APR-2002.
 XX
 XX 03-OCT-2001; 2001WO-US30821.
 PF
 XX 03-OCT-2000; 2000US-237189P.
 PR
 XX
 XX (GENE-) GENE LOGIC INC.
 PA
 XX Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
 PI WPI: 2002-435328/46.
 XX
 DR
 XX
 PT Detecting granulocyte activation by detecting differential expression
 PT of genes associated with granulocyte activation, which serves as
 PT diagnostic markers that is useful for monitoring disease states and
 PT drug toxicity
 XX
 XX Claim 1; SEQ ID NO 137; 114pp; English.
 XX
 XX The invention relates to detecting (M1) granulocyte (GC) activation
 CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by
 CC DNA chip analysis as given in the specification, and comparing
 CC the expression level to an expression level in an unactivated
 CC GC, where differential expression of Gs is indicative of GCA.
 CC Also included are modulating (M2) GA by contacting GC with an agent
 CC that alters the expression of at least one gene in Gs; (2) screening (M3)
 CC for an agent capable of modulating GCA or an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease using the
 CC gene expression profile; (3) detecting (M4) an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease, by detecting the
 CC level of expression in a sample of the tissue of gene(s) from Gs, where
 CC the level of expression of the gene is indicative of inflammation;
 CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,
 CC an allergic response in a subject, exposure of a subject to a pathogen
 CC or sterile inflammatory disease, by contacting a tissue having
 CC inflammation with an agent that modulates the expression of gene(s)
 CC from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for
 CC modulating GCA; M3 is useful for screening an agent capable of modulating
 CC GCA preferably in an inflammation in a tissue; M4 is useful for
 CC detecting an inflammation (especially chronic) in a tissue, an allergic
 CC response in a subject, exposure of a subject to a pathogen or sterile
 CC inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
 CC glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
 CC reperfusion injury, ARDS, adult respiratory distress syndrome,
 CC inflammatory bowel disease, Crohn's disease, ulcerative colitis,
 CC periodontal disease; also bacterial infection, viral infection,
 CC parasitic infection, protozoal infection, fungal infection and M5 is
 CC useful for treating one of the above conditions. The present
 CC sequence represents a gene differentially expressed in granulocytes.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pcl_sequences.
 XX
 SQ Sequence 147724 BP; 46968 A; 29251 C; 28325 G; 43180 T; 0 other;
 Query Match 5.6%; Score 22; DB 24; Length 147724;
 Best Local Similarity 100.0%; Pred. No. 0.66;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 149 GGCCCTGTGTCATGCGCTCAC 170
 ||||||||||||||||||
 Db 123545 GGCCCTGTGTCATGCGCTCAC 123524
 RESULT 7

AAS91061/c
 ID AAS91061 standard; cDNA: 1179 BP.
 AC AAS91061;
 XX
 DT 13-FEB-2002 (first entry)
 DE DNA encoding novel human diagnostic protein #2685.
 DE Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 OS
 PN WO200175067-A2.
 PD 11-OCT-2001.
 PF 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 XX 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 PI WPI: 2001-639362/73.
 DR P-PSDB: ABC26874.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS Claim 1; SEQ ID No 2685; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 1179 BP; 348 A; 292 C; 315 G; 224 T; 0 other;
 Query Match 5.1%; Score 20; DB 23; Length 1179;
 Best Local Similarity 100.0%; Pred. No. 6.7;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 191 GGGTGCAGATCTGTGGG 210
 ||||||||||||||||||||
 DB 284 GGGTGCAGATCTGTGGG 265
 RESULT 8
 AAS66674
 ID AAS66674 standard; cDNA: 2526 BP.

XX
 AC AAS66674;
 XX
 DT 13-FEB-2002 (first entry)
 DE DNA encoding novel human diagnostic protein #2478.
 DE Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 OS
 PN WO200175067-A2.
 PD 11-OCT-2001.
 PF 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 XX 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 PI WPI: 2001-639362/73.
 DR P-PSDB: ABC02487.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS Claim 1; SEQ ID No 2478; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 2526 BP; 871 A; 536 C; 541 G; 578 T; 0 other;
 Query Match 5.1%; Score 20; DB 23; Length 2526;
 Best Local Similarity 100.0%; Pred. No. 6.6;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 32 TTTCATCTTTCAGGACTT 51
 ||||||||||||||||||||
 DB 2377 TTTCATCTTTCAGGACTT 2396
 RESULT 9
 AAC90702/c
 ID AAC90702 standard; cDNA: 366 BP.

```

XX 14-MAR-2001 (first entry)
DT
XX
DE Human secretory protein TSC-456 nucleotide sequence SEQ ID NO:17.
XX
KM Human; secretory protein; cancer; immune disease; infectious disease;
KM lung function disorder; liver function disorder; antiinflammatory;
KM gastrointestinal disorder; cytostatic; haematopoietic; anticoagulant;
KM immunomodulatory; hepatotropic; cell proliferation-stimulant;
KM cell migratory agent; cell differentiation-inducer; ss.
XX
OS Homo sapiens.
XX
PN WO200071581-A1.
XX
PD 30-NOV-2000.
XX
PE 19-MAY-2000; 2000WO-JP03221.
XX
XX 20-MAY-1999; 99JP-0140229.
XX
XX (TAKE ) TAKEDA CHEM IND LTD.
XX
XX Itoh Y, Mogi S, Tanaka H, Ohkubo S, Ogi K,
XX WPI: 2001-032023/04.
XX DR P-PSDB; AAB36662.
XX
XX Novel secretory protein and its salt with e.g. anti-cancer,
XX anti-inflammatory and hematopoietic, effects, applicable as drugs in
XX remedies and preventives to treat diseases like cancer and immune
XX diseases
XX
XX Example 2; Page 97-98; 122pp; Japanese.
XX
XX AAC90701 to AAC90715 encode the human secretory proteins given in
XX AAB36661 to AAB36675. The proteins can have cytostatic,
XX anti-inflammatory, haematopoietic, anti-coagulant, immunomodulatory and
XX hepatotropic activities, and can be used as cell migratory agents, cell
XX proliferation-stimulants and cell differentiation-inducers. The proteins
XX are useful in the treatment and prevention of diseases such as cancer,
XX lung function disorder, liver function disorder, gastrointestinal
XX disorder and immune diseases. AAC90716 to AAC90755 represent PCR primers
XX which are used in the exemplification of the present invention.
XX
XX Sequence 366 BP; 99 A; 97 C; 83 G; 87 T; 0 other;
XX
XX Query Match 4.8%; Score 19; DB 22; Length 366;
XX Best Local Similarity 100.0%; Pred. No. 21;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 217 GTGCTTCTCGATGGGGT 235
DB 238 GTGCTTCTCGATGGGGT 220

```

```

KM endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.
XX
XX Homo sapiens.
XX
XX WO9924836-A1.
XX
XX 20-MAY-1999.
XX
XX 04-NOV-1998; 98WO-US23435.
XX
XX 17-NOV-1997; 97US-0066100.
XX PR 07-NOV-1997; 97US-0064900.
XX PR 07-NOV-1997; 97US-0064908.
XX PR 07-NOV-1997; 97US-0064911.
XX PR 07-NOV-1997; 97US-0064912.
XX PR 07-NOV-1997; 97US-0064913.
XX PR 07-NOV-1997; 97US-0064984.
XX PR 07-NOV-1997; 97US-0064985.
XX PR 07-NOV-1997; 97US-0064987.
XX PR 17-NOV-1997; 97US-0066090.
XX PR 17-NOV-1997; 97US-0066094.
XX PR 17-NOV-1997; 97US-0066095.
XX PR 17-NOV-1997; 97US-0066089.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Carter KC, Ebner R, Endress GA, Feng P, Janat F;
XX PI Ryan H, Laileur DW, Moore PA, Ni J, Olsen HS, Rosen CA;
XX PI Ruben SM, Shi Y, Soppet DR, Wei Y;
XX DR WPI: 1999-337740/28.
XX DR P-PSDB; AAY27583.
XX
XX New human secreted proteins and coding sequences useful for treating
XX disorders of the immune system and hyperproliferative disorders
XX
XX Claim 1; Page 290; 507pp; English.
XX
XX This sequence represents a nucleic acid molecule which encodes a
XX secreted human protein. The gene number is given in the descriptor line.
XX The gene can be used to generate fusion proteins by linking to the gene
XX to a human immunoglobulin Fc portion (e.g. AAX84924) for increasing the
XX stability of the fused protein as compared to the human protein only.
XX The invention relates to 125 novel genes and their fragments (nucleic
XX acid sequences: AAX84933-X85057; amino acid sequences AAY27567-Y27933)
XX which are useful for preventing, treating or ameliorating medical
XX conditions e.g. by protein or gene therapy. Also, pathological
XX conditions can be diagnosed by determining the amount of the new
XX polypeptides in a sample or by determining the presence of mutations in
XX the new polynucleotides. Specific uses are described for each of the 125
XX polynucleotides, based on which tissues they are most highly expressed in
XX (see AAX84933 for described uses).
XX
XX Sequence 802 BP; 208 A; 210 C; 190 G; 188 T; 6 other;
XX
XX Query Match 4.8%; Score 19; DB 20; Length 802;
XX Best Local Similarity 100.0%; Pred. No. 21;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 217 GTGCTTCTCGATGGGGT 235
DB 304 GTGCTTCTCGATGGGGT 286

```

```

RESULT 11
AAC68089/C
ID AAC68089 standard; CDNA; 1982 BP.
XX
AC AAC68089;
XX
XX 20-FEB-2001 (first entry)
DT
XX

```

```

Human; secreted protein; fusion protein; gene therapy; protein therapy;
diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;
developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;

```


DE Human secreted protein cDNA sequence #9.
KW Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral;
KW anti-allelic; hepatotropic; antidiabetic; anti-inflammatory; anticancer;
KW vulnere; anticonvulsant; antibacterial; antifungal; antiparasitic;
KW candid; gene therapy; cancer; immune disorder; cardiovascular disorder;
KW neurological disease; infection; human; secreted protein; ss.
XX
OS Homo sapiens.
XX
XX MO200058335-A1.
XX
PD 05-OCT-2000.
XX
PF 22-MAR-2000; 2000WO-US07534.
XX
PR 26-MAR-1999; 99US-0126598.
PR 22-DEC-1999; 99US-0171504.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA (ROSE/) ROSEN C A.
PI Rosen CA, Ruben SM, Komatsoulis G;
XX
XX WPI: 2000-611702/58.
DR P-PSDB; AAB37356.
XX
PT Nucleic acids encoding human secreted proteins, used to treat, prevent,
PT ameliorate or diagnose conditions such as cancer, and autoimmune
PT diseases e.g. arthritis -
XX
XX
PS Claim 1; Pages 321-322; 387pp; English.
XX
CC The invention relates to the isolation of genes AAC68081-C68127 encoding
CC 47 human secreted proteins AAB37348-B37394. The genes can be used to
CC generate fusion proteins by linking to the gene for the human
CC immunoglobulin G Fc portion (AAC68072) for increasing the stability of
CC the fusion protein as compared to the human protein only. The genes and
CC proteins are useful for preventing, ameliorating or treating medical
CC conditions, e.g. by protein or gene therapy. The genes are isolated
CC from a range of human tissues disclosed in the specification. The
CC nucleic acids, proteins, antibodies and (ant)agonists are useful in
CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
CC and ovarian cancer, and other cancers of the adrenal gland, bone, bone
CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
CC colitis; (c) cardiovascular disorders such as myocardial ischaemias; (d)
CC wound healing; (e) neurological diseases e.g. cerebral anoxia and
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
CC and parasitic infections.
XX
XX
SQ Sequence 1982 BP; 549 A; 513 C; 471 G; 448 T; 1 other;
XX
XX
XX Query Match 4.8%; Score 19; DB 21; Length 1982;
XX Best Local Similarity 100.0%; Pred. NO. 21;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0.
XX
OY 326 TTTTCTCATTTCTCTGTGG 344
XX |TTTTTTTTTTTTTTTT|
DB 860 TTTTCTCATTTCTCTGTGG 842
XX
RESULT 12
ABLO9112
ID ABLO9112 standard; cDNA; 2770 BP.
XX
XX ABL09112;
AC
DT 26-MAR-2002 (first entry)
XX
DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 21819.

XX	Drosophila; developmental biology; cell signalling; insecticide;
KW	Drosophila; developmental biology; cell signalling; insecticide;
KW	pharmaceutical; gene; ss.
OS	Drosophila melanogaster.
XX	WO200171042-A2.
XX	27-SEP-2001.
XX	
PF	23-MAR-2001; 2001WO-US09231.
XX	
XX	23-MAR-2000; 2000US-191637P.
PR	11-JUL-2000; 2000US-0614150.
XX	
PA	(PEKE) PE CORP NY.
XX	
FI	Venter JC, Adams M, Li PWD, Myers EW;
XX	
DR	WPI; 2001-656860/75.
DR	P-PSDB; ABB65009.
PT	New isolated nucleic acid detection reagent for detecting 1000 or more
PT	genes from Drosophila and for elucidating cell signalling and cell-cell
PT	interactions -
XX	
PS	Claim 1; SEQ ID NO 21818; 21pp + Sequence Listing; English.
XX	
CC	The invention relates to an isolated nucleic acid detection reagent
CC	capable of detecting 1000 or more genes from Drosophila. The invention is
CC	useful in developmental biology and in elucidating cell signalling and
CC	cell-cell interactions in higher eukaryotes for the development of
CC	insecticides, therapeutics and pharmaceutical drugs. The invention
CC	discloses genomic DNA sequences (ABL01840-ABL16175), expressed DNA
CC	sequences (ABL01840-ABL16175) and the encoded proteins
CC	(ABB57737-ABB72072).
CC	The sequence data for this patent did not form part of the printed
CC	specification, but was obtained in electronic format directly from WIPO
CC	at ftp.wipo.int/pub/published_pct_sequences.
XX	
XX	Sequence 2770 BP; 786 A; 625 C; 629 G; 730 T; 0 other;
XX	
Query Match	4.8%; Score 19; DB 23; Length 2770;
Best Local Similarity	100.0%; Pred. No. 20;
Matches 19; Conservative	0; Mismatches 0; Indels 0; Gaps 0
OY	233 GGTCAACAATCACTCACT 251
Db	1050 GGTCAACAATCACTCACT 1068
XX	
RESULT 13	
ABL28930/c	
ID	ABL28930 standard; DNA; 4902 BP.
XX	
AC	ABL28930;
XX	
DT	26-MAR-2002 (first entry)
XX	
DE	Drosophila melanogaster genomic polynucleotide SEQ ID NO 36263.
XX	
KW	Drosophila; developmental biology; cell signalling; insecticide;
KW	pharmaceutical; gene; ds.
XX	
OS	Drosophila melanogaster.
XX	
PN	WO200171042-A2.
XX	
PD	27-SEP-2001.
XX	
PF	23-MAR-2001; 2001WO-US09231.
XX	
PR	23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.
XX
XX (PEKE) PE CORP NY.
XX
PI Venter JC, Adams M, Li PWD, Myers EW.
XX
XX WPI: 2001-656860/75.
XX
XX New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from Drosophila and for elucidating cell signalling and cell-cell
PT interactions -
XX
XX Claim 1; SEQ ID NO 38263; 21pp + Sequence Listing; English.
XX
XX The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from Drosophila. The invention is
CC useful in developmental biology and in elucidating cell signalling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA
CC sequences (AB116176-AB116175) and the encoded proteins
CC (AB116177-AB116178).
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 4902 BP; 1509 A; 1006 C; 984 G; 1403 T; 0 other;
Query Match 4.8%; Score 19; DB 23; Length 4902;
Best Local Similarity 100.0%; Pred. No. 20;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 268 GGTGGGCTTTTGTTCG 286
D 886 GGTGGGCTTTTGTTCG 868
RESULT 14
AA145064
ID AA145064 standard; DNA; 31 BP.
XX
XX AA145064;
AC
XX 05-FEB-1997 (first entry)
DT
XX
DE 21-hydroxylase fragment A, forward primer.
XX
XX Polymerase chain reaction; primer; PCR; amplify; 21-hydroxylase gene;
human adrenal gland cDNA library; diagnosis; detection;
Addison's disease; ss.
XX
XX Syntheitic.
OS
XX
XX USS376533-A.
PN
XX
XX 27-DEC-1994.
PD
XX
XX 24-MAY-1993; 93US-0066281.
PE
XX
XX 24-MAY-1993; 93US-0066281.
PR
XX
XX (UYFL) UNIV FLORIDA.
PA
XX
XX MacIaren NK, Song YH;
PI
XX
XX WPI: 1995-043462/06.
DR
XX
XX Detection of Addison's disease or persons at risk from developing it
PT - using a 21-hydroxylase peptide fragment to detect autoantibodies
PT associated with Addison's disease.
XX
XX Disclosure; Column 6; 9pp; English.
XX

CC The sequences given in AA145064-75 are primers which were used to
CC amplify fragments of the 21-hydroxylase gene from a human adrenal
CC gland cDNA library. The amplified fragments encode peptides which
CC were used in the method of the invention to diagnose or detect
CC Addison's disease. Fragment A comprises amino acids 1-162, fragment
CC B, amino acids 164-356, fragment C, amino acids 3440494, fragment D,
CC amino acids 164-271, fragment E, amino acids 272-356, and fragment F
CC amino acids 197-298 of the 21-hydroxylase enzyme. These primers
CC have been optimised for use and do not directly correspond to 21
CC hydroxylase gene sequences. All the forward primers contain BamHI
CC sites and all the reverse primers contain EcoRI restriction sites.
XX
SQ Sequence 31 BP; 2 A; 10 C; 11 G; 8 T; 0 other;
Query Match 4.5%; Score 18; DB 16; Length 31;
Best Local Similarity 100.0%; Pred. No. 67;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 288 TCCATGCTGCTCTGCGC 305
D 8 TCCATGCTGCTCTGCGC 25
RESULT 15
AA243080/C
ID AA243080 standard; DNA; 339 BP.
XX
XX AA243080;
AC
XX
XX 04-FEB-2000 (first entry)
DT
XX
DE C. elegans insulin-like protein ZK84.6 gene.
XX
XX Insulin-like protein; diagnosis; insulin-like gene analysis; nematode;
KW insulin hormone; aging; senescence; pesticide target; signalling pathway;
KW signal transduction pathway; ss.
XX
XX Caenorhabditis elegans.
OS
XX
XX W09954436-A2.
PN
XX
XX 28-OCT-1999.
PD
XX
XX 16-APR-1999; 99WO-US08522.
PE
XX
XX 17-APR-1998; 98US-0062580.
PR
XX 08-MAY-1998; 98US-0074984.
PR 26-MAY-1998; 98US-0084303.
XX
XX (EXEL-) EXELIXIS PHARM INC.
PA
XX
XX Homburger SA, Platt DM, Ferguson KC, Doberstein SK, Buchman AR;
PI Reddy BP;
PI
XX
XX WPI: 2000-012239/01.
DR
XX
XX P-PSDB; AAY65657.
DR
XX
XX Analysing Caenorhabditis elegans insulin-like gene expression, nucleic
PT acids and proteins of the C. elegans insulin-like genes -
PT
XX
XX Claim 4; Fig 8; 194pp; English.
PS
XX
XX This sequence encodes a Caenorhabditis elegans insulin-like protein,
CC and can be used in the method of the invention. The method is for
CC analysing an effect of expression or mis-expression of a C. elegans
CC insulin-like gene, and comprises observing a first nematode genetically
CC engineered to express or mis-express a C. elegans insulin-like protein
CC (ILP) of any one of groups I, II or IV or a derivative or fragment that
CC displays one or more functional activities of the C. elegans ILP. The
CC insulin-like genes in C. elegans constitute very useful tools for probing
CC the function and regulation of their corresponding pathways. This can be
CC expected to lead to the discovery of new drug targets, therapeutic
CC proteins, diagnostics and prognostics useful in the treatment of diseases

CC and clinical problems associated with the function of insulin hormones in
 CC humans and other animals, as well as clinical problems associated with
 CC aging and senescence. The information may also be useful in
 CC identification and validation of pesticide targets in invertebrate pests
 CC that are components of these signalling pathways. The genes are also
 CC useful for identifying factors that are upstream of the receptor in the
 CC signal transduction pathway. The ligand-encoding C. elegans insulin-like
 CC genes provide a superior approach for identifying factors that are
 CC upstream of the receptor in the signal transduction pathway.

CC
 XX Sequence 339 BP; 92 A; 86 C; 72 G; 89 T; 0 other;

Query Match 4.5%; Score 18; DB 21; Length 339;

Best local Similarity 100.0%; Pred. No. 65;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 188 TGTGGTTGCCAAGATCT 205

|||||

Db 260 TGTGGTTGCCAAGATCT 243

Search completed: April 25, 2003, 00:01:03
 Job time : 220.988 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:49:04 ; Search time 1311.72 Seconds
(without alignments)
4815.901 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atggctgagcttccttcgcctt.....gamctgatacttcagtga 396

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 16154066 seqs, 8097743376 residues

d size : 0

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estmu:*
5: em_estrov:*
6: em_estrpl:*
7: em_estro:*
8: em_hlc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rtd:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	147	37.1	470	17	A0770688 HS_5368_B
2	146	36.9	525	17	A0165256 HS_3025_B
3	38	9.6	628	17	A0237815 RPCI11-70
4	35	8.8	410	17	A0442274 HS_5137_A
5	35	8.8	453	17	A0437684 HS_5137_A
6	35	8.8	635	17	A0390599 CITBI-El-

7	34	8.6	412	17	A0321916 RPCI11-10	
c	34	8.6	529	17	A0881246 HS_5137_B	
c	9	378	17	A0479650 RPCI-11-2	A0479650 RPCI-11-2	
c	10	8.3	471	17	A0147593 HS_3065_B	A0147593 HS_3065_B
c	11	8.3	482	17	A0320567 RPCI11-59	A0320567 RPCI11-59
c	12	8.3	546	12	BF962702 PM4-NM120	BF962702 PM4-NM120
c	13	8.3	563	17	A0420187 RPCI-11-1	A0420187 RPCI-11-1
c	14	8.3	691	17	A2516454 RPCI-11-1	A2516454 RPCI-11-1
c	15	8.3	723	17	A0386439 RPCI11-15	A0386439 RPCI11-15
c	16	8.1	427	17	A0697116 HS_5528_A	A0697116 HS_5528_A
c	17	7.8	553	17	A2521751 RPCI-11-1	A2521751 RPCI-11-1
c	18	7.8	695	17	AG179297 Pan trogl	AG179297 Pan trogl
c	19	7.8	769	17	A0899390 HS_5234_A	A0899390 HS_5234_A
c	20	7.6	360	17	A0207172 HS_3238_B	A0207172 HS_3238_B
c	21	7.6	399	17	A0115544 RPCI11-57	A0115544 RPCI11-57
c	22	7.6	435	17	A0116061 RPCI11-57	A0116061 RPCI11-57
c	23	7.6	452	17	A0442744 HS_5122_B	A0442744 HS_5122_B
c	24	7.6	551	17	A0369689 HS_5333_B	A0369689 HS_5333_B
c	25	7.3	414	17	A0003326 RPCI11-25	A0003326 RPCI11-25
c	26	7.3	468	17	A0819715 HS_5513_A	A0819715 HS_5513_A
c	27	7.3	615	17	AG161224 Pan trogl	AG161224 Pan trogl
c	28	7.3	653	17	AG143347 Pan trogl	AG143347 Pan trogl
c	29	7.1	363	17	A0120796 HS_3076_A	A0120796 HS_3076_A
c	30	7.1	376	17	A0548294 RPCI-11-4	A0548294 RPCI-11-4
c	31	7.1	401	17	A0588089 CITBI-El-	A0588089 CITBI-El-
c	32	7.1	419	17	A0436209 HS_5049_B	A0436209 HS_5049_B
c	33	7.1	425	17	A0141070 HS_3141_B	A0141070 HS_3141_B
c	34	7.1	449	17	A0269217 RPCI11-69	A0269217 RPCI11-69
c	35	7.1	480	17	A0817757 HS_5250_B	A0817757 HS_5250_B
c	36	7.1	530	17	A0193128 HS_3060_B	A0193128 HS_3060_B
c	37	7.1	541	17	A0683783 HS_5455_B	A0683783 HS_5455_B
c	38	7.1	653	17	AG160919 Pan trogl	AG160919 Pan trogl
c	39	7.1	681	17	AG141287 Pan trogl	AG141287 Pan trogl
c	40	7.1	684	17	AG049745 Pan trogl	AG049745 Pan trogl
c	41	6.8	345	17	A0075574 CIT-HSP-2	A0075574 CIT-HSP-2
c	42	6.8	517	17	A0614252 HS_5123_B	A0614252 HS_5123_B
c	43	6.8	529	17	A0346593 RPCI11-12	A0346593 RPCI11-12
c	44	6.8	553	17	A0238365 RPCI11-63	A0238365 RPCI11-63
c	45	6.8	606	17	A0350708 RPCI11-11	A0350708 RPCI11-11

ALIGNMENTS

RESULT 1
LOCUS A0770688
DEFINITION HS_5368_B2.C08.SP66 RPCI-11 Human Male BAC Library Homo sapiens
ACCESSION A0770688
VERSION A0770688.1 GI:5648804
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 470)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong

(pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm) or from Research Genetics (<http://www.hsc.washington.edu>)
http://www.hsc.washington.edu
Plate: 944 Row: F Column: 16
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 470.

FEATURES
source
Location/Qualifiers
1..470

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=944 Col=16 Row=F"
/clone_11b="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"

BASE COUNT 83 a 112 c 131 g 141 t 3 others

Query Match 37.1%; Score 147; DB 17; Length 470;
Best Local Similarity 100.0%; Pred. No. 4.1e-54;
Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGCTTGGGCGG 60
|||||
Db 103 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGCTTGGGCGG 162

QY 61 GAGTATGTAACACTCTGCTCTGTGTGTGCTGAGTGCTGCTACTGAGACTCTG 120
|||||
Db 163 GAGTATGTAACACTCTGCTCTGTGTGTGCTGAGTGCTGCTACTGAGACTCTG 222

QY 121 CATACACAGCTCTGATATGAGACCA 147
|||||
Db 223 CATACACAGCTCTGATATGAGACCA 249

RESULT 2 525 bp DNA linear GSS 16-OCT-1998
AO165256
LOCUS
DEFINITION HS-3025-B2-G06-T7 CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone Plate=3025 Col=12 Row=N, DNA sequence.
ACCESSION AO165256
VERSION AO165256.1 GI:3563451
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 525)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3025 Row: N Column: 12
Class: BAC ends
High quality sequence stop: 525.

FEATURES
source
Location/Qualifiers
1..525

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3025 Col=12 Row=N"
/clone_11b="CIT Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC clones in
E-Coli DH10B"

BASE COUNT 102 a 139 c 137 g 143 t 4 others

Query Match 36.9%; Score 146; DB 17; Length 525;
Best Local Similarity 99.5%; Pred. No. 1.1e-53;
Matches 196; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 80 GTCTCTGTGTGCTCTGATGCTGCTCTACGTAGAGACTCTGCATACAGCTCTGATAT 139
|||||
Db 145 GTCTCTGTGTGCTCTGATGCTGCTCTACGTAGAGACTCTGCATACAGCTCTGATAT 204

QY 140 CGGACCCAGGCGCTGTGTCATGAGGCTCAGAGAGATCCCGTATCTGTGGTTGCA 199
|||||
Db 205 CGGACCCAGGCGCTGTGTCATGAGGCTCAGAGAGATCCCGTATCTGTGGTTGCA 264

QY 200 AGATCTGTGGAGAACTGTGTCTCTGATGAGGCTCAGACATCAGCTCTTCCCT 259
|||||
Db 265 AGATCTGTGGAGAACTGTGTCTCTGATGAGGCTCAGACATCAGCTCTTCCCT 324

QY 260 TGGCTGAGCTGGGCT 276
|||||
Db 325 TGGCTGAGCTGGGCT 341

RESULT 3 628 bp DNA linear GSS 21-APR-1999
AO237815
LOCUS
DEFINITION RPC111-70H4.TK RPCI-11 Homo sapiens genomic clone RPCI-11-70H4, DNA
sequence.
ACCESSION AO237815
VERSION AO237815.1 GI:3670106
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 628)
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdamas@tigr.org

Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genetics (http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html)
Seq primer: 77
Class: BAC ends.

FEATURES
source
Location/Qualifiers
1..628

/organism="Homo sapiens"
/db_xref="GDB:752667"
/db_xref="taxon:9606"
/clone="RPCI-11-70H4"
/clone_11b="RPCI-11"
/sex="male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

BASE COUNT 125 a 150 c 164 g 189 t
ORIGIN
Query Match 9.6%; Score 38; DB 17; Length 628;
Best Local Similarity 100.0%; Pred. No. 2.6e-06;
Matches 38; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 61 GAGTATGTAACCTCCGGCTCTGTGTGCTGAG 98
|||||
Db 157 GAGTATGTAACCTCCGGCTCTGTGTGCTGAG 194
|||||
RESULT 4 410 bp DNA linear GSS 31-MAR-1999
LOCUS A0442274
DEFINITION HS_5137_A1.P12.SP6E RPCT-11 Human Male BAC Library Homo sapiens
genomic clone Plate=713 Col=23 Row=K, DNA sequence.
ACCESSION A0442274
VERSION A0442274.1 GI:4553613
MEDLINE
JOURNAL
COMMENT human
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 410)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
LOCUS are derived from the human BAC library RPCT-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: K column: 23
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 410.
FEATURES
Location/Qualifiers
1. 410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=23 Row=K"
/clone_lib="RPCT-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"
BASE COUNT 80 a 111 c 111 g 107 t
ORIGIN
Query Match 8.8%; Score 35; DB 17; Length 410;
Best Local Similarity 100.0%; Pred. No. 6e-05;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 108
|||||
Db 160 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 194
|||||

RESULT 5 453 bp DNA linear GSS 31-MAR-1999
LOCUS A0437684
DEFINITION HS_5137_A2.H06.SP6E RPCT-11 Human Male BAC Library Homo sapiens
genomic clone Plate=713 Col=12 Row=O, DNA sequence.
ACCESSION A0437684
VERSION A0437684.1 GI:4549023
MEDLINE
JOURNAL
COMMENT human
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 453)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
LOCUS are derived from the human BAC library RPCT-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: O column: 12
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 453.
FEATURES
Location/Qualifiers
1. 453
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=12 Row=O"
/clone_lib="RPCT-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"
BASE COUNT 84 a 127 c 117 g 124 t
ORIGIN
Query Match 8.8%; Score 35; DB 17; Length 453;
Best Local Similarity 100.0%; Pred. No. 5.8e-05;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 74 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 108
|||||
Db 163 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 197
|||||
RESULT 6 635 bp DNA linear GSS 06-MAR-1999
LOCUS A0390599/c
DEFINITION CITBI-El-2544B15.TR CITBI-El Homo sapiens genomic clone 2544B15,
DNA sequence.
ACCESSION A0390599
VERSION A0390599.1 GI:4361622
MEDLINE
JOURNAL
COMMENT human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS	Zhao,S., Adams,M.D., Nierman,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.
TITLE	Use of BAC End Sequences from Caltech Libraries for Sequence-Ready Map Building
JOURNAL	Unpublished (1997)
COMMENT	Other.GSS: CITBI-EI-2544BI5.TF Contact: Shaying Zhao, William Nierman, Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbeetlgr.org Clones are available from Research Genetics (info@resgen.com). BAC end search page: http://www.tlgr.org/tldb/hungen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends.
URES	Location/Qualifiers
source	1..635 /organism="Homo sapiens" /db_xref="taxon:9606" /clone="2544BI5" /clone_1lb="CITBI-EI" /sex="male" /cell_type="sperm" /note="Vector: pBeloBAC11; Site_1: EcoRI; Site_2: EcoRI; Caltech Human BAC Library P"
BASE COUNT	192 a 144 c 172 g 127 t
ORIGIN	
Query Match	8.8%; Score 35; DB 17; Length 635;
Best Local Similarity	100.0%; Pred. No. 5,4e-05;
Matches	35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db	527 TCCTGGCTCTGTGTGTGCTGAGTGGCTCTCTC 108 TCCTGGCTCTGTGTGTGCTGAGTGGCTCTCTC 493
RESULT 7	
LOCUS	AQ321916 412 bp DNA linear GSS 06-MAY-1999
DEFINITION	RPC11-101H18.TV RPC1-11 Homo sapiens genomic clone RPC1-11-101H18, DNA sequence.
ACCESSION	AQ321916
VERSION	AQ321916.1 GI:4054584
KEYWORDS	GSS.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS	1 (bases 1 to 412) Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Liber,K., Golden,K., Berry,K., Granger,D., Sub,E., White,C., de Jong,P. and Venter,J.C.
TITLE	Use of human BAC End Sequences for Sequence-Ready Map Building
JOURNAL	Unpublished (1998)
COMMENT	Other.GSS: RPC11-101H18.TV Contact: Shaying Zhao, William Nierman, Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbeetlgr.org Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Plier de Jong (plierdejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tlgr.org/tldb/hungen/bac_end_search/bac_end_search.html

```

FEATURES
  source
    location/Qualifiers
      1..412
        /organism="Homo sapiens"
        /db_xref="GDB:7538585"
        /db_xref="taxon:9606"
        /clone="RPC1-11-101H18"
        /clone_1fb="RPC1-11"
        /sex="Male"
        /cell_type="Lymphocytes"
        /note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
        Rcc11 Human Male BAC library"
BASE COUNT      74 a      102 c      120 g      116 t
ORIGIN
Query Match      8.6%; Score 34; DB 17; Length 412;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 191 GGGTTGCAAGATCTGTGGAGACAGATGTGGTTTC 224
    |||||||
Db 292 GGGTTGCAAGATCTGTGGAGACAGATGTGGTTTC 325

```

[illegible]

BASE COUNT 137 a 147 c 121 g 117 t 7 others
ORIGIN pBAC3.6 vector at EcoRI sites"

Query Match 8.6%; Score 34; DB 17; Length 529;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAAACCTCGGCTCTGTGTTCG 93
DB 382 GGAGTATGTAAACCTCGGCTCTGTGTTCG 349

RESULT 9
LOCUS A0479650/c 378 bp DNA linear GSS 23-APR-1999
DEFINITION RPCI-11-269D19.TV RPCI-11 Homo sapiens genomic clone RPCI-11-269D19
ACCESSION A0479650
VERSION A0479650.1 GI:4661769
WORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 378)
Zhaio,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter
J.C.
TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
JOURNAL Unpublished (1997)
CONTACT: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeetlgr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
Research Genet. cs (info@resgen.com). BAC end search page:
http://www.tlgr.org/tldb/hungun/bac_end_search/bac_end_search.html.
Seq primer: 17
Class: BAC ends.

FEATURES
Location/Qualifiers
1..378
/organism="Homo sapiens"
/db_xref="GDB:7603002"
/db_xref="taxon:9606"
/clone="RPCI-11-269D19"
/clone_lib="RPCI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC Library"

BASE COUNT 116 a 103 c 93 g 66 t

Query Match 8.3%; Score 33; DB 17; Length 378;
Best Local Similarity 100.0%; Pred. No. 0.00046;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTCAAGATCTGTGGAGAGTGTTCG 225
DB 208 GTTCAAGATCTGTGGAGAGTGTTCG 176

RESULT 10
LOCUS A0147593/c 471 bp DNA linear GSS 08-OCT-1998
DEFINITION HS_3065_B2_H08_MF CIT Approved Human Genomic Sperm Library D Homo

ACCESSION sapiens genomic clone Plate=3065 Col=16 Row=P, DNA sequence.
VERSION A0147593
KEYWORDS A0147593.1 GI:3538246
SOURCE GSS.
ORGANISM human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 471)
Mahiras,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahiras GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3065 Row: P Column: 16
Class: BAC ends
High quality sequence stop: 471.
Location/Qualifiers
1..471
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3065 Col=16 Row=P"
/clone_lib="CIT Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT 144 a 118 c 123 g 84 t 2 others

ORIGIN

Query Match 8.3%; Score 33; DB 17; Length 471;
Best Local Similarity 100.0%; Pred. No. 0.00044;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTCAAGATCTGTGGAGAGTGTTCG 225
DB 265 GTTCAAGATCTGTGGAGAGTGTTCG 233

RESULT 11
LOCUS A0320567/c 482 bp DNA linear GSS 04-MAY-1999
DEFINITION RPCI11-99N1.TV RPCI-11 Homo sapiens genomic clone RPCI-11-99N1, DNA
sequence.
ACCESSION A0320567
VERSION A0320567.1 GI:4050696
WORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 482)
Adams,M.D., Rounsley,S.D., Zhao,S., Baas,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
TITLE Use of human BAC End Sequences for Sequence-Ready Map Building
JOURNAL Unpublished (1998)
COMMENT Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeetlgr.org
Clones are derived from the human BAC library RPCI-11. For BAC

library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html
Seq primer: 17
Class: BAC ends.

FEATURES

Location/Qualifiers

1..482

/organism="Homo sapiens"

/db_xref="GDB:7537944"

/db_xref="taxon:9606"

/clone="RPCI-11-99N1"

/clone_1id="RPCI-11"

/sex="Male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

RPC111 Human Male BAC Library"

BASE COUNT 142 a 121 c 106 g 112 t 1 others

ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 0.00044;

Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 53 TGGGCGCGAGTATGTAACCTCGGCTCT 85
|||||
Db 338 TGGGCGCGAGTATGTAACCTCGGCTCT 306

RESULT 12 546 bp mRNA linear EST 22-JAN-2001
LOCUS BF962702
DEFINITION PM4-NM1209-151200-012-f04 NM1209 Homo sapiens cDNA, mRNA sequence.
ACCESSION BF962702
VERSION BF962702.1 GI:12379977
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 546)
Dias Neto E., Garcia Correa R., Verjovski-Almeida S., Briones M.R.,
Nagal M.A., da Silva M. Jr., Zago M.A., Bordin S., Costa F.F.,
Goldman G.H., Carvalho A.F., Matsukuma A., Bala G.S., Simpson D.H.,
Brunstein A., de Oliveira P.S., Bucher P., Jongeneel C.V., O'Hare
M.J., Soares F., Brentani R.R., Reis L.F., de Souza S.J. and
Simpson A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663

JOURNAL COMMENT
MEDLINE
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel.: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?cl=PM4&t2=PM4-NM1209-151200-012-f04&t3=2000-12-15&t4=1>)
Seq primer: puc 18 forward
High quality sequence start: 15
High quality sequence stop: 545.

FEATURES

Location/Qualifiers

1..546

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone_1id="NM1209"

/dev_stage="Adult"

/note="Organ: nervous system; Vector: puc18; Site_1: SmaI;
Site_2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

BASE COUNT 149 a 157 c 135 g 105 t

ORIGIN

Query Match 8.3%; Score 33; DB 12; Length 546;

Best Local Similarity 100.0%; Pred. No. 0.00044;

Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTGCAAGATCTGTGGAGAGTGTGTTCC 225
|||||
Db 96 GTTGCAAGATCTGTGGAGAGTGTGTTCC 64

RESULT 13 563 bp DNA linear GSS 23-MAR-1999
LOCUS AQ420187
DEFINITION RPCI-11-185J19-TV RPCI-11 Homo sapiens genomic clone RPCI-11-185J19
DNA sequence.
ACCESSION AQ420187
VERSION AQ420187.1 GI:4477911
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 563)
Zhao S., Adams M.D., Nierman W., Malek J., de Jong P. and Venter
J.C.
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
Unpublished (1997)
Other GSSs: RPCI-11-185J19-TV
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genet cs (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html
Seq primer: SP6
Class: BAC ends.

FEATURES Location/Qualifiers

source

1..563

/organism="Homo sapiens"

/db_xref="GDB:7570890"

/db_xref="taxon:9606"

/clone="RPCI-11-185J19"

/clone_1id="RPCI-11"

/sex="Male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

RPC111 Human Male BAC Library"

BASE COUNT 173 a 127 c 114 g 149 t

ORIGIN

Query Match 8.3%; Score 33; DB 17; Length 563;

Best Local Similarity 100.0%; Pred. No. 0.00042;

Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 53 TGGGCGCGAGTATGTAACCTCGGCTCT 85
|||||

DB 333 TGGGGCCGAGATATGTAACCTCTGGTCTCT 301

RESULT 14
LOCUS AZ516454/c
DEFINITION RPCI-11-191D19.TV RPCI-11 Homo sapiens genomic clone RPCI-11-191D19
' DNA sequence.
ACCESSION AZ516454
VERSION AZ516454.1 GI:10824878
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Eukaryota; Eutheria; Primates; Catarrhini; Homnidae; Homo.
Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
J.C.
BAC end sequences of library RPCI-11
Unpublished (1997)
Other_GSSs: RPCI-11-191D19.TVB
Contact: Shaying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
BACPAC Resources (http://bacpac.med.buffalo.edu). Clones may be purchased from
Research Genet. cs (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.
This BAC end was generated during the RAD process and may have
higher chance of clone tracking errors.
Seq primer: T7
Class: BAC ends.

FEATURES
Location/Qualifiers
1..691
/organism="Homo sapiens"
/db_xref="GDB:7573050"
/db_xref="taxon:9606"
/clone="RPCI-11-191D19"
/clone_1lb="RPCI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC library"

BASE COUNT 203 a 191 c 174 g 121 t 2 others

ORIGIN

Query Match 8.3%; Score 33; DB 17; Length 691;
Best Local Similarity 100.0%; Pred. No. 0.0004;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTGCAAGATCTGTGGAGAGTGTGTTCC 225
|||||
DB 216 GTTGCAAGATCTGTGGAGAGTGTGTTCC 184

RESULT 15
LOCUS AO386439/c
DEFINITION RPCI11-154D6.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-154D6,
DNA sequence.
ACCESSION AO386439
VERSION AO386439.1 GI:4357462
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Eukaryota; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 723)
AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
J.C.
TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
JOURNAL Unpublished (1997)
COMMENT Other_GSSs: RPCI11-154D6.TV
Contact: Shaying Zhao, William Niernan, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeet@tigr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
Research Genet. cs (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html
Seq primer: SP6
Class: BAC ends.

FEATURES
Location/Qualifiers
1..723
/organism="Homo sapiens"
/db_xref="GDB:7558829"
/db_xref="taxon:9606"
/clone="RPCI-11-154D6"
/clone_1lb="RPCI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC library"

BASE COUNT 224 a 165 c 135 g 199 t

ORIGIN

Query Match 8.3%; Score 33; DB 17; Length 723;
Best Local Similarity 100.0%; Pred. No. 0.0004;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 53 TGGGGCCGAGATATGTAACCTCTGGGCTCT 85
|||||
DB 331 TGGGGCCGAGATATGTAACCTCTGGGCTCT 299

Search completed: April 25, 2003, 00:52:51
Job time : 1334.72 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:57:29 ; Search time 46.8639 Seconds

(without alignments)
2591.419 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_446

Perfect score: 396
Sequence: 1 atgggtgacatcttcgcctt.....gamctgatacttcagtga 396

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 441362 seqs, 153338381 residues

d size : 0

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : Issued_Patents_NA:*

1: /cgn2_6/ptodata/1/ina/5A_COMB.seq:*
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq:*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq:*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq:*
5: /cgn2_6/ptodata/1/ina/PCITUS_COMB.seq:*
6: /cgn2_6/ptodata/1/ina/Backfillseq1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	4.5	31	1	US-08-066-281-1
2	18	4.5	430	4	US-09-397-787-254
3	17	4.3	1118	4	US-09-452-239-37
4	17	4.3	1146	4	US-09-452-239-3
5	17	4.3	1381	4	US-08-858-207A-108
6	17	4.3	1509	1	US-08-115-052-1
7	17	4.3	2686	4	US-09-228-986-3
8	17	4.3	11613	4	US-09-453-702B-42
9	17	4.3	40328	3	US-08-742-185-102
10	16	4.0	36	4	US-08-910-722-7
11	16	4.0	42	4	US-08-910-722-5
12	16	4.0	57	1	US-08-474-177-17
13	16	4.0	57	1	US-08-487-033-17
14	16	4.0	57	1	US-08-480-810-17
15	16	4.0	57	2	US-08-508-735-17
16	16	4.0	57	2	US-08-848-251-17
17	16	4.0	57	2	US-08-848-047-17
18	16	4.0	57	3	US-09-120-130-17
19	16	4.0	57	3	US-09-115-252-17
20	16	4.0	57	3	US-08-986-515-17
21	16	4.0	57	3	US-09-120-128-17
22	16	4.0	57	4	US-09-120-129-17
23	16	4.0	57	4	US-09-201-139-17
24	16	4.0	57	4	US-09-120-131-17
25	16	4.0	384	4	US-09-134-001C-1752
26	16	4.0	471	1	US-08-474-177-1
27	16	4.0	471	1	US-08-487-033-1

C 28	16	4.0	471	1	US-08-480-810-1	Sequence 1, Appli
C 29	16	4.0	471	2	US-08-508-735-1	Sequence 1, Appli
C 30	16	4.0	471	2	US-08-848-251-1	Sequence 1, Appli
C 31	16	4.0	471	2	US-08-486-047-1	Sequence 1, Appli
C 32	16	4.0	471	3	US-09-120-130-1	Sequence 1, Appli
C 33	16	4.0	471	3	US-09-115-252-1	Sequence 1, Appli
C 34	16	4.0	471	3	US-08-986-515-1	Sequence 1, Appli
C 35	16	4.0	471	3	US-09-120-128-1	Sequence 1, Appli
C 36	16	4.0	471	4	US-09-120-129-1	Sequence 1, Appli
C 37	16	4.0	471	4	US-09-201-139-1	Sequence 1, Appli
C 38	16	4.0	471	4	US-09-120-131-1	Sequence 1, Appli
C 39	16	4.0	536	4	US-09-221-017B-1081	Sequence 1081, Ap
C 40	16	4.0	687	4	US-09-457-568-23	Sequence 23, Appl
C 41	16	4.0	687	4	US-09-457-646-23	Sequence 23, Appl
C 42	16	4.0	737	4	US-09-457-568-19	Sequence 19, Appl
C 43	16	4.0	737	4	US-09-457-646-19	Sequence 19, Appl
C 44	16	4.0	782	4	US-09-457-568-21	Sequence 21, Appl
C 45	16	4.0	782	4	US-09-457-646-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1
US-08-066-281-1
Sequence 1, Application US/08066281
Patent No. 5376533
GENERAL INFORMATION:
APPLICANT: MacIaren, No. 53765331 K.
ATTORNEY/AGENT INFORMATION:
TITLE OF INVENTION: Methods and Compositions for the Detection of
TITLE OF INVENTION: Addison's Disease
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESSES:
ADDRESSEE: DAVID R. SALIWANCHIK
STREET: 2421 N.W. 41st Street, Suite A-1
CITY: Gainesville
STATE: FL
COUNTRY: USA
ZIP: 32606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/066,281
FILING DATE: 19930524
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: SalIWanchik, David R.
REGISTRATION NUMBER: 31,794
REFERENCE/DOCKET NUMBER: UF/S&S-132
TELECOMMUNICATION INFORMATION:
TELEPHONE: 904-375-8100
TELEFAX: 904-372-5800
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 31 bases
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (synthetic)
US-08-066-281-1
Query Match 4.5%; Score 18; DB 1; Length 31;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 288 TCCATGCTGCTCTGAGGC 305
|||||
Db 8 TCCATGCTGCTCTGAGGC 25

```
RESULT 2
US-09-397-787-254/C
: Sequence 254, Application US/09397787
: Patent No. 6468758
: GENERAL INFORMATION:
: APPLICANT: Benson, Darin R.
: APPLICANT: Lodes, Michael J.
: APPLICANT: Mitcham, Jennifer L.
: APPLICANT: King, Gordon E.
: TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
: FILE REFERENCE: 210121.466C2
: CURRENT APPLICATION NUMBER: US/09/397,787
: NUMBER OF SEQ ID NOS: 334
: SOFTWARE: FastSeq for Windows Version 3.0
: SEQ ID NO 254
: LENGTH: 430
: TYPE: DNA
: ORGANISM: Homo sapien
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)..(430)
: OTHER INFORMATION: n = A,T,C or G
US-09-397-787-254

Query Match          4.3%; Score 18; DB 4; Length 430;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 191 GCGTTGCAAGATCTGCG 208
DB 273 GCGTTGCAAGATCTGCG 256

RESULT 3
US-09-452-239-37
: Sequence 37, Application US/09452239
: Patent No. 6465229
: GENERAL INFORMATION:
: APPLICANT: Rafalski, Antoni J.
: APPLICANT: Fader, Gary M.
: APPLICANT: Cahoon, Rebecca E.
: TITLE OF INVENTION: Plant Caffey1-CoA O-Methyltransferase
: FILE REFERENCE: BB1284 US NA
: CURRENT APPLICATION NUMBER: US/09/452,239
: EARLIER FILING DATE: 1999-12-01
: EARLIER APPLICATION NUMBER: 60/110,594
: NUMBER OF SEQ ID NOS: 50
: SOFTWARE: Microsoft Office 97
: SEQ ID NO 37
: LENGTH: 1118
: TYPE: DNA
: ORGANISM: Trifolium aestivum
US-09-452-239-37

Query Match          4.3%; Score 17; DB 4; Length 1118;
Best Local Similarity 100.0%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 273 GCGTTTTTTTGGCTC 289
DB 971 GCGTTTTTTTGGCTC 987

RESULT 4
US-09-452-239-3
: Sequence 3, Application US/09452239
: Patent No. 6465229
: GENERAL INFORMATION:
: APPLICANT: Rafalski, Antoni J.

US-08-858-207A-108/C
: Sequence 108, Application US/08858207A
: Patent No. 6348328
: GENERAL INFORMATION:
: APPLICANT: Black, Michael
: APPLICANT: Hodgson, John
: APPLICANT: Knowles, David
: APPLICANT: Nicholas, Richard
: APPLICANT: Stodola, Robert
: TITLE OF INVENTION: NO. 6348328e1 Compounds
: NUMBER OF SEQUENCES: 552
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SmithKline Beecham Corporation
: STREET: 709 Swedeland Road
: CITY: King of Prussia
: STATE: PA
: COUNTRY: USA
: ZIP: 19406-0939
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette
: COMPUTER: IBM Compatible
: OPERATING SYSTEM: DOS
: SOFTWARE: FastSeq for Windows Version 2.0
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/858,207A
: FILING DATE: 09-MAY-1997
: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: 60/017670
: FILING DATE: 14-MAY-1996
: ATTORNEY/AGENT INFORMATION:
: NAME: Gimml, Edward R
: REGISTRATION NUMBER: 38,891
: REFERENCE/DOCKET NUMBER: P50475
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 610-270-4478
: TELEFAX: 610-270-5090
: TELEX:
: INFORMATION FOR SEQ ID NO: 108:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1381 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-08-858-207A-108

Query Match          4.3%; Score 17; DB 4; Length 1381;
```


TITLE OF INVENTION: No. 6365723el Sequences of E. coli 0157
NUMBER OF SEQUENCES: 265
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Quarles & Brady
STREET: 1 South Pinckney Street
CITY: Madison
STATE: WI
COUNTRY: US
ZIP: 53701-2113
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44mb storage
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Word Perfect 8.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/453, 702B
FILING DATE: 03-Dec-1999
CLASSIFICATION: <unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/110,955
FILING DATE: 04-DEC-1998
ATTORNEY/AGENT INFORMATION:
NAME: Seay, Nicholas J.
REGISTRATION NUMBER: 27386
REFERENCE/DOCKET NUMBER: 960296.95017
TELECOMMUNICATION INFORMATION:
TELEPHONE: (608) 251-5000
TELEFAX: (608) 251-9166
INFORMATION FOR SEQ ID NO: 42:
SEQUENCE CHARACTERISTICS:
LENGTH: 11613
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 42:
US-09-453-702B-42

Query Match 4.3%; Score 17; DB 4; Length 11613;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 219 GGTTCCTGATGGGCT 235
|||||
DB 2106 GGTTCCTGATGGGCT 2090

RESULT 9
Sequence 102, Application US/08742185
Patent No. 6020476

GENERAL INFORMATION:
APPLICANT: Page, David C.
APPLICANT: Reljo, Renee
APPLICANT: Saxena, Richa
APPLICANT: Hawkins, Trevor
APPLICANT: Reene, Mary Pat
TITLE OF INVENTION: DAZ: A GENE FAMILY ASSOCIATED WITH AZOOSPERMIA
NUMBER OF SEQUENCES: 102
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Millita Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: US
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/742,185

FILING DATE: 30-OCT-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/690,734
FILING DATE: 31-JUL-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/310,429
FILING DATE: 22-SEP-1994
ATTORNEY/AGENT INFORMATION:
NAME: Granahan, Patricia
REGISTRATION NUMBER: 32,227
REFERENCE/DOCKET NUMBER: WH194-07A2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 861-6240
TELEFAX: (617) 861-9540
INFORMATION FOR SEQ ID NO: 102:
SEQUENCE CHARACTERISTICS:
LENGTH: 40328 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-742-185-102

Query Match 4.3%; Score 17; DB 3; Length 40328;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 268 GGTGGGGCTTTT TTTT 284
|||||
DB 882 GGTGGGGCTTTT TTTT 898

RESULT 10
US-08-910-722-7/c
Sequence 77, Application US/08910722
Patent No. 6251871

GENERAL INFORMATION:
APPLICANT: Jin, Xiaomei
APPLICANT: Roth, Jack A.
TITLE OF INVENTION: p16 EXPRESSION CONSTRUCTS AND THEIR
TITLE OF INVENTION: APPLICATION IN CANCER THERAPY
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Arnold, White & Durkee
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: United States of America
ZIP: 77210
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,722
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/502,881
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Highlander, Steven L.
REGISTRATION NUMBER: 37,642
REFERENCE/DOCKET NUMBER: INGN-016/HYL
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
TELE: 79-0924
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs

TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "DNA"
US-08-910-722-7

Query Match 4.0%; Score 16; DB 4; Length 36;
Best Local Similarity 100.0%; Pred. No. 93;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 285 GGCTCCATGCTGCTCC 300
|||||
DB 23 GGCTCCATGCTGCTCC 8

RESULT 11
US-08-910-722-5/C
Sequence 5, Application US/08910722
Patent No. 6251871

GENERAL INFORMATION:

APPLICANT: Jin, Xiaomei
APPLICANT: Roth, Jack A.
TITLE OF INVENTION: p16 EXPRESSION CONSTRUCTS AND THEIR
TITLE OF INVENTION: APPLICATION IN CANCER THERAPY
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Arnold, White & Durkee
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: United States of America
ZIP: 77210
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,722
FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/502,881
FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Highlander, Steven L.
REGISTRATION NUMBER: 37,642
REFERENCE/DOCKET NUMBER: INGN:016/HYL

TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577

TELEX: 79-0924

INFORMATION FOR SEQ ID NO: 5:

SEQUENCE CHARACTERISTICS:
LENGTH: 42 base pairs
TYPE: nucleic acid

STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "DNA"

US-08-910-722-5

Query Match 4.0%; Score 16; DB 4; Length 42;
Best Local Similarity 100.0%; Pred. No. 93;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 285 GGCTCCATGCTGCTCC 300
|||||
DB 29 GGCTCCATGCTGCTCC 14

RESULT 12

US-08-474-177-17/C
Sequence 17, Application US/08474177
Patent No. 5624819

GENERAL INFORMATION:

APPLICANT: Skolnick, Mark H.
APPLICANT: Cannon-Albright, Lisa A.
TITLE OF INVENTION: GERMLINE MUTATIONS IN THE MTS GENE
NUMBER OF SEQUENCES: 36
CORRESPONDENCE ADDRESS:

ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
STREET: 1201 New York Avenue, Suite 1000
CITY: Washington
STATE: DC

COUNTRY: USA
ZIP: 20005

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/474,177
FILING DATE: 07-JUN-1995

CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/03537
FILING DATE: 17-MAR-1995

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/251,938
FILING DATE: 01-JUN-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/215,087
FILING DATE: 18-MAR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/215,086
FILING DATE: 18-MAR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/227,369
FILING DATE: 14-APR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/214,582
FILING DATE: 18-MAR-1994

ATTORNEY/AGENT INFORMATION:

NAME: Ihnen, Jeffrey L.
REGISTRATION NUMBER: 28,957
REFERENCE/DOCKET NUMBER: 24884-109348-E

TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-962-4810

TELEFAX: 202-962-8300

INFORMATION FOR SEQ ID NO: 17:

SEQUENCE CHARACTERISTICS:
LENGTH: 57 base pairs
TYPE: nucleic acid

STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO

ANTI-SENSE: NO
ORIGINAL SOURCE:

ORGANISM: Homo sapiens

US-08-474-177-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 285 GGCTCCATGCTGCTCC 300
|||||
DB 50 GGCTCCATGCTGCTCC 35

RESULT 13

US-08-487-033-17/c
; Sequence 17, Application US/08487033
; Patent No. 5739027
; GENERAL INFORMATION:
; APPLICANT: Kamb, Alexander
; TITLE OF INVENTION: MTS1-Beta GENE
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
; STREET: 1201 New York Avenue, Suite 1000
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/487,033
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03316
; FILING DATE: 17-MAR-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/251,938
; FILING DATE: 01-JUN-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,087
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,086
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/227,369
; FILING DATE: 14-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/214,582
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ihnen, Jeffrey L.
; REGISTRATION NUMBER: 28,957
; REFERENCE/DOCKET NUMBER: 24884-109348-C
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-962-8300
; TELEFAX: 202-962-8300
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 57 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-08-487-033-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 285 GGCTCCATGCTGCTCC 300
|||||
Db 50 GGCTCCATGCTGCTCC 35

RESULT 14
US-08-480-810-17/c
; Sequence 17, Application US/08480810

; Patent No. 5801236
; GENERAL INFORMATION:
; APPLICANT: Kamb, Alexander
; TITLE OF INVENTION: MTS1 GENE
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
; STREET: 1201 New York Avenue, Suite 1000
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/480,810
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03316
; FILING DATE: 17-MAR-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/251,938
; FILING DATE: 01-JUN-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,087
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,086
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/227,369
; FILING DATE: 14-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/214,582
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ihnen, Jeffrey L.
; REGISTRATION NUMBER: 28,957
; REFERENCE/DOCKET NUMBER: 24884-109348
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-962-8300
; TELEFAX: 202-962-8300
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 57 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-08-480-810-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 285 GGCTCCATGCTGCTCC 300
|||||
Db 50 GGCTCCATGCTGCTCC 35

RESULT 15
US-08-508-735-17/c
; Sequence 17, Application US/08508735
; Patent No. 5843756
; GENERAL INFORMATION:

APPLICANT: Stone, Steven
 APPLICANT: Jang, Ping
 APPLICANT: Kamb, Alexander
 TITLE OF INVENTION: MTS GENE AND THERAPEUTIC USE THEREOF
 NUMBER OF SEQUENCES: 47
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
 STREET: 1201 New York Avenue, Suite 1000
 CITY: Washington
 STATE: DC
 COUNTRY: USA
 ZIP: 20005
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/508,735
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US to be assigned
 FILING DATE: 07-JUN-1995
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: PCT/US95/03316
 FILING DATE: 17-MAR-1995
 ATTORNEY/AGENT INFORMATION:
 NAME: Ihnen, Jeffrey L.
 REGISTRATION NUMBER: 28,957
 REFERENCE/DOCKET NUMBER: 24884-109348
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-962-8300
 TELEFAX: 202-962-4848
 INFORMATION FOR SEQ ID NO: 17:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 57 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 ORIGINAL SOURCE:
 ORGANISM: Homo sapiens
 US-08-508-735-17

Query Match 4.0%; Score 16; DB 2; Length 57;
 Best Local Similarity 100.0%; Pred. NO. 92;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 285 GGCTCCATGCTGCTCC 300
 ||||||||||||
 Db 50 GGCTCCATGCTGCTCC 35

Search completed: April 25, 2003, 00:54:16
 Job time : 56.8639 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:41:49 ; Search time 81.2308 Seconds
(without alignments)
5304.628 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atgggtgacatttcgctt.....gamctgatacttcagtga 396

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 709820 seqs, 544064369 residues

d size: 0

Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

Published_Applications_NA:*

- 1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/ptodata/1/pubpna/PC1_NEW_PUB.seq:*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
- 6: /cgn2_6/ptodata/1/pubpna/PC1US_PUBCOMB.seq:*
- 7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
- 8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
- 9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
- 10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
- 12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
- 13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
- 14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	33	8.3	684973	10	US-09-263-959-1
C 2	22	5.6	361	9	US-09-918-995-8549
C 3	19	4.8	802	9	US-09-974-879-27
C 4	18	4.5	430	10	US-09-876-889-254
C 5	18	4.5	431	10	US-09-866-562-80
C 6	18	4.5	431	10	US-09-866-562-87
C 7	18	4.5	570	10	US-09-864-761-9118
C 8	17	4.3	489	9	US-09-918-995-28049
C 9	17	4.3	491	9	US-09-918-995-34650
C 10	17	4.3	531	9	US-09-918-995-19262
C 11	17	4.3	531	9	US-10-092-154-1899
C 12	17	4.3	553	10	US-09-764-847-1899
C 13	17	4.3	553	10	US-09-864-761-13668
C 14	17	4.3	609	10	US-09-974-300-8084
C 15	17	4.3	944	9	US-09-774-639-99
C 16	17	4.3	944	9	US-09-969-730-16
C 17	17	4.3	1047	10	US-09-823-830A-468
C 18	17	4.3	1089	10	US-09-962-740-3
C 19	17	4.3	1118	10	US-09-452-239-37

20	17	4.3	1146	10	US-09-452-239-3	Sequence 3, Appli
C 21	17	4.3	1215	10	US-09-962-740-1	Sequence 1, Appli
C 22	17	4.3	1215	10	US-09-962-740-6	Sequence 6, Appli
C 23	17	4.3	1396	10	US-09-962-740-8	Sequence 8, Appli
C 24	17	4.3	1631	10	US-09-962-740-5	Sequence 5, Appli
C 25	17	4.3	2686	9	US-10-101-464A-3	Sequence 3, Appli
C 26	17	4.3	2880	9	US-09-951-502A-1	Sequence 1, Appli
C 27	17	4.3	11613	9	US-10-114-170-42	Sequence 42, Appli
C 28	17	4.3	38374	10	US-09-880-107-3463	Sequence 3463, Ap
C 29	17	4.3	172637	10	US-09-805-458A-3	Sequence 3, Appli
C 30	17	4.3	326014	10	US-09-731-231A-3	Sequence 1, Appli
C 31	17	4.3	1503841	9	US-09-946-807-1	Sequence 1, Appli
C 32	17	4.3	1503841	10	US-09-795-668-1	Sequence 1, Appli
C 33	17	4.3	1503841	10	US-09-795-668-1	Sequence 1, Appli
C 34	16	4.0	121	9	US-09-818-875-1321	Sequence 1321, Ap
C 35	16	4.0	121	9	US-09-818-875-1322	Sequence 1322, Ap
C 36	16	4.0	121	9	US-09-818-875-1325	Sequence 1325, Ap
C 37	16	4.0	121	9	US-09-818-875-1326	Sequence 1326, Ap
C 38	16	4.0	121	9	US-09-818-875-1329	Sequence 1329, Ap
C 39	16	4.0	121	9	US-09-818-875-1330	Sequence 1330, Ap
C 40	16	4.0	121	9	US-09-818-875-1333	Sequence 1333, Ap
C 41	16	4.0	121	9	US-09-818-875-1334	Sequence 1334, Ap
C 42	16	4.0	238	10	US-09-878-574-5259	Sequence 5259, Ap
C 43	16	4.0	235	10	US-09-878-574-13480	Sequence 13480, A
C 44	16	4.0	281	10	US-09-964-824A-385	Sequence 385, App
C 45	16	4.0	321	10	US-09-864-761-25530	Sequence 25530, A

ALIGNMENTS

RESULT 1

US-09-263-959-1/c

Sequence 1, Application US/09263959

Patent No. US20020150891A1

GENERAL INFORMATION:

APPLICANT: Hood, Leroy E.

APPLICANT: Koop, Ben F.

TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U

NUMBER OF SEQUENCES: 1279

CORRESPONDENCE ADDRESS:

ADDRESSEE: Seed and Berry LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue

CITY: Seattle

STATE: Washington

COUNTRY: US

ZIP: 98104-7092

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/263,959

FILING DATE: 05-MAR-1999

CLASSIFICATION:

ATTORNEY/AGENT INFORMATION:

NAME: Mcmasters, David D.

REGISTRATION NUMBER: 33,963

REFERENCE/DOCKET NUMBER: 920010.426C2

TELECOMMUNICATION INFORMATION:

TELEPHONE: (206) 622-4900

TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 684973 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-09-263-959-1

Query Match 8.3% Score 33; DB 10; Length 684973;

Best Local Similarity 100.0%; Pred. No. 6,4e-08;
Matches 33: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 66 TGTAAACTCCGGCTCTGTGTGCTGAG 98
|||||

Db 404690 TGTAAACTCCGGCTCTGTGTGCTGAG 404658

RESULT 2

US-09-918-995-8549/c

; Sequence 8549, Application US/09918995
; Publication No. US20030073623A1

; GENERAL INFORMATION:

; APPLICANT: Hyseq, Inc.

; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED

; FILE REFERENCE: 20411-756

; CURRENT APPLICATION NUMBER: US/09/918,995

; PRIOR FILING DATE: 2001-07-30

; PRIOR APPLICATION NUMBER: US/09/235,076

; NUMBER OF SEQ ID NOS: 38054

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 8549

; LENGTH: 361

; TYPE: DNA

; ORGANISM: Homo sapiens

US-09-918-995-8549

Query Match Best Local Similarity 100.0%; Pred. No. 0.063;
Matches 22: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 149 GGCCTGTGTCATGGCTC 170
|||||

Db 143 GGCCTGTGTCATGGCTC 122

RESULT 3

US-09-974-879-27/c

; Sequence 27, Application US/09974879

; Publication No. US20030028003A1

; GENERAL INFORMATION:

; APPLICANT: Rosen et al.

; TITLE OF INVENTION: 125 Human Secreted Proteins

; FILE REFERENCE: P2020P2

; CURRENT APPLICATION NUMBER: US/09/974,879

; PRIOR FILING DATE: 2001-10-12

; PRIOR APPLICATION NUMBER: US 60/239,893

; PRIOR FILING DATE: 2000-10-13

; PRIOR APPLICATION NUMBER: US 09/818,683

; PRIOR FILING DATE: 2001-03-28

; PRIOR APPLICATION NUMBER: US 09/305,736

; PRIOR FILING DATE: 1999-05-05

; PRIOR APPLICATION NUMBER: PCT/US98/23435

; PRIOR FILING DATE: 1998-11-04

; PRIOR APPLICATION NUMBER: US 60/064,911

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,912

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,983

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,900

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,988

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,987

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,908

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,984

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/064,985

; PRIOR FILING DATE: 1997-11-07

; PRIOR APPLICATION NUMBER: US 60/066,094

; PRIOR FILING DATE: 1997-11-17

; PRIOR APPLICATION NUMBER: US 60/066,100

; PRIOR FILING DATE: 1997-11-17

; PRIOR APPLICATION NUMBER: US 60/066,089

; PRIOR FILING DATE: 1997-11-17

; PRIOR APPLICATION NUMBER: US 60/066,095

; PRIOR FILING DATE: 1997-11-17

; PRIOR APPLICATION NUMBER: US 60/066,090

; PRIOR FILING DATE: 1997-11-17

; NUMBER OF SEQ ID NOS: 611

; SOFTWARE: PatentIn Ver. 2.0

; SEQ ID NO 27

; LENGTH: 802

; TYPE: DNA

; ORGANISM: Homo sapiens

; FEATURE:

; NAME/KEY: SITE

; LOCATION: (337)

; OTHER INFORMATION: n equals a,t,g, or c

; NAME/KEY: SITE

; LOCATION: (359)

; OTHER INFORMATION: n equals a,t,g, or c

US-09-974-879-27

Query Match Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 19: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 217 GTGGTTCTGATGGGT 235
|||||

Db 304 GTGGTTCTGATGGGT 286

RESULT 4

US-09-876-889-254/c

; Sequence 254, Application US/09876889

; Patent No. US20020076715A1

; GENERAL INFORMATION:

; APPLICANT: Benson, Darin R.

; APPLICANT: Lodes, Michael J.

; APPLICANT: Mitcham, Jennifer L.

; APPLICANT: King, Gordon E.

; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN

; FILE REFERENCE: 210121.466C3

; CURRENT APPLICATION NUMBER: US/09/876,889

; NUMBER OF SEQ ID NOS: 353

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 254

; LENGTH: 430

; TYPE: DNA

; ORGANISM: Homo sapien

; FEATURE:

; NAME/KEY: misc.feature

; LOCATION: (1) --(430)

; OTHER INFORMATION: n = A,T,C or G

US-09-876-889-254

Query Match Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 18: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 191 GGGTGCAGATCTGTG 208
|||||

Db 273 GGGTGCAGATCTGTG 256

RESULT 5

US-09-866-562-80

; Sequence 80, Application US/0986562

```

Patent No. US20020009758A1
GENERAL INFORMATION:
APPLICANT: Harlocker, Susan L.
APPLICANT: Wang, Tongtong
APPLICANT: Bangur, Chaitanya S.
APPLICANT: Klee, Jennifer
APPLICANT: Switzer, Anne
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
FILE REFERENCE: 210121.502
CURRENT APPLICATION NUMBER: US/09/866,562
CURRENT FILING DATE: 2001-05-25
NUMBER OF SEQ ID NOS: 96
SEQ ID NO 80
LENGTH: 431
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc-feature
LOCATION: 361,431
OTHER INFORMATION: n = A,T,C or G
09-866-562-80

```

```

Query Match 4.5%; Score 18; DB 10; Length 431;
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 91 TGCTGAGTGGCTGCTCT 108
DB 18 TGCTGAGTGGCTGCTCT 35

```

```

RESULT 6
US-09-866-562-87
Sequence 87, Application US/09866562
Patent No. US20020009758A1
GENERAL INFORMATION:
APPLICANT: Harlocker, Susan L.
APPLICANT: Wang, Tongtong
APPLICANT: Bangur, Chaitanya S.
APPLICANT: Klee, Jennifer
APPLICANT: Switzer, Anne
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
FILE REFERENCE: 210121.502
CURRENT APPLICATION NUMBER: US/09/866,562
CURRENT FILING DATE: 2001-05-25
NUMBER OF SEQ ID NOS: 96
SEQ ID NO 87
LENGTH: 431
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc-feature
LOCATION: 361,431
OTHER INFORMATION: n = A,T,C or G
US-09-866-562-87

```

```

Query Match 4.5%; Score 18; DB 10; Length 431;
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 91 TGCTGAGTGGCTGCTCT 108
DB 18 TGCTGAGTGGCTGCTCT 35

```

```

RESULT 7
US-09-864-761-9118/c
Sequence 9118, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.

```

```

APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO
FILE REFERENCE: Aecm1ca-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
SEQ ID NO 9118
LENGTH: 570
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000053.1
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.8
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 2.4
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 2.3
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
US-09-864-761-9118

```

```

Query Match 4.5%; Score 18; DB 10; Length 570;
Best Local Similarity 100.0%; Pred. No. 8.6;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 85 TGTGTGCTGAGTGGC 102
DB 505 TGTGTGCTGAGTGGC 488

```

```

RESULT 8
US-09-918-995-28049/c

```

```
; Sequence 28049, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 28049
; LENGTH: 489
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(489)
; OTHER INFORMATION: n = A,T,C or G
; US-918-995-28049
```

```
Query Match 4.3%; Score 17; DB 9; Length 489;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 325 TTTTCTTCATCTCTG 341
|||||
DB 416 TTTTCTTCATCTCTG 400
```

```
RESULT 9
US-09-918-995-34650/C
; Sequence 34650, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 34650
; LENGTH: 491
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(491)
; OTHER INFORMATION: n = A,T,C or G
; US-09-918-995-34650
```

```
Query Match 4.3%; Score 17; DB 9; Length 491;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 328 TTTCTTCATCTCTGTG 344
|||||
DB 138 TTTCTTCATCTCTGTG 122
```

```
RESULT 10
US-09-918-995-19262/C
; Sequence 19262, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
```

```
; TITLE OF INVENTION: FROM VARIOUS CDNA LIBRARIES
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19262
; LENGTH: 492
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(492)
; OTHER INFORMATION: n = A,T,C or G
; US-09-918-995-19262
```

```
Query Match 4.3%; Score 17; DB 9; Length 492;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 246 CTCACGCTTCCTCTGG 262
|||||
DB 85 CTCACGCTTCCTCTGG 69
```

```
RESULT 11
US-10-092-154-1899/C
; Sequence 1899, Application US/10092154
; Publication No. US20030054575A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009C1
; CURRENT APPLICATION NUMBER: US/10/092,154
; CURRENT FILING DATE: 2002-03-07
; NUMBER OF SEQ ID NOS: 2003
; PRIOR Application removed - See File Wrapper or Palm
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-092-154-1899
```

```
Query Match 4.3%; Score 17; DB 9; Length 531;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 28 TCTTTTCATCTTGCA 44
|||||
DB 141 TCTTTTCATCTTGCA 125
```

```
RESULT 12
US-09-764-847-1899/C
; Sequence 1899, Application US/09764847
; Patent No. US20020132767A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009
; CURRENT APPLICATION NUMBER: US/09/764,847
; CURRENT FILING DATE: 2001-01-17
; PRIOR application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 2003
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-764-847-1899
```


Query Match 4.3%; Score 17; DB 10; Length 531;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 28 TCCTTTTCATCTTGCA 44
 Db 141 TCCTTTTCATCTTGCA 125

RESULT 13
 US-09-864-761-13668/c
 ; Sequence 13668, Application US/09864761
 ; Patent No. US2002048763A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Penn, Sharon G.
 ; APPLICANT: Rank, David R.
 ; APPLICANT: Hanzel, David K.
 ; APPLICANT: Chen, Wensheng
 ; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
 ; FILE REFERENCE: Aecomica-X-1
 ; CURRENT APPLICATION NUMBER: US/09/864,761
 ; PRIOR FILING DATE: 2001-05-23
 ; PRIOR APPLICATION NUMBER: US 60/180,312
 ; PRIOR FILING DATE: 2000-02-04
 ; PRIOR APPLICATION NUMBER: US 60/207,456
 ; PRIOR FILING DATE: 2000-05-26
 ; PRIOR APPLICATION NUMBER: US 09/632,366
 ; PRIOR FILING DATE: 2000-08-03
 ; PRIOR APPLICATION NUMBER: GB 24263.6
 ; PRIOR FILING DATE: 2000-10-04
 ; PRIOR APPLICATION NUMBER: US 60/236,359
 ; PRIOR FILING DATE: 2000-09-27
 ; PRIOR APPLICATION NUMBER: PCT/US01/00666
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00667
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00664
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00669
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00665
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00668
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00663
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00662
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00661
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00670
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: US 60/234,687
 ; PRIOR FILING DATE: 2000-09-21
 ; PRIOR APPLICATION NUMBER: US 09/608,408
 ; PRIOR FILING DATE: 2000-06-30
 ; PRIOR APPLICATION NUMBER: US 09/774,203
 ; PRIOR FILING DATE: 2001-01-29
 ; NUMBER OF SEQ ID NOS: 49117
 ; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
 ; SEQ ID NO 13668
 ; LENGTH: 553
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; OTHER INFORMATION: MAP TO 282201.1
 ; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.4
 US-09-864-761-13668

Query Match 4.3%; Score 17; DB 10; Length 553;
 Best Local Similarity 100.0%; Pred. No. 30; Mismatches 0; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 327 TTCTTCATCTCTGCG 343
 Db 378 TTCTTCATCTCTGCG 362

RESULT 14
 US-09-974-300-8084/c
 ; Sequence 8084, Application US/09974300
 ; Patent No. US20020146721A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Berka, Randy M.
 ; APPLICANT: Clausen, Ib Groth
 ; TITLE OF INVENTION: Methods For Monitoring Multiple Gene
 ; FILE REFERENCE: 10085.500-US
 ; CURRENT APPLICATION NUMBER: US/09/974,300
 ; PRIOR FILING DATE: 2001-10-05
 ; PRIOR APPLICATION NUMBER: 09/680,598
 ; PRIOR FILING DATE: 2000-10-06
 ; PRIOR APPLICATION NUMBER: 60/279,526
 ; PRIOR FILING DATE: 2001-03-27
 ; NUMBER OF SEQ ID NOS: 8481
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 8084
 ; LENGTH: 609
 ; TYPE: DNA
 ; ORGANISM: Bacillus clausii
 US-09-974-300-8084

Query Match 4.3%; Score 17; DB 10; Length 609;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCCTTTTCATCTTGCA 44
 Db 536 TCCTTTTCATCTTGCA 520

RESULT 15
 US-09-774-639-99/c
 ; Sequence 99, Application US/09774639
 ; Publication No. US2003000355A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Rosen et al.
 ; TITLE OF INVENTION: 90 Human Secreted Proteins
 ; FILE REFERENCE: P2013P1
 ; CURRENT APPLICATION NUMBER: US/09/774,639
 ; PRIOR FILING DATE: 2001-07-09
 ; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/244,112
 ; PRIOR FILING DATE: EARLIER FILING DATE: 1999-02-04
 ; NUMBER OF SEQ ID NOS: 371
 ; SOFTWARE: Patentln Ver. 2.0
 ; SEQ ID NO 99
 ; LENGTH: 944
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; NAME/KEY: SITE
 ; LOCATION: (13)
 ; OTHER INFORMATION: n equals a,t,g, or c
 ; NAME/KEY: SITE
 ; LOCATION: (486)
 ; OTHER INFORMATION: n equals a,t,g, or c
 ; NAME/KEY: SITE
 ; LOCATION: (934)
 ; OTHER INFORMATION: n equals a,t,g, or c
 US-09-774-639-99

Query Match 4.3%; Score 17; DB 9; Length 944;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 246 CTCACCTGCTTCCTGG 262
|||||
Db 515 CTCACCTGCTTCCTGG 499

Search completed: April 25, 2003, 02:12:39
Job time : 286.231 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: April 24, 2003, 22:30:38 ; Search time 1361 Seconds
(without alignments)
791.186 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSEFALODSFSSLOGLLGPEYVKLLGLCVCLSGCST 37

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

ched: 2054640 segs, 14551402878 residues
Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODE=frame+ p2n.model -DEV=xlh
-Q/cgn2.1/USPTO_spool/US09513999/unat_18042003.170936-28357/app-query.fasta.1.199
-DB=GenEmbl -QFMT=fastap -SUFFIX=p2n.rge -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=-1 -MATRIX=biosum62 -TRANS=human40.cdi -LIST=45
-DOCALLIGN=200 -THR.SCORE=pct -THR.MAX=100 -THR.MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=fto -NCM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn.1.1.1687_etunat.18042003.170936.28357 -NCPU=6 -ICPU=3
-NO_XUPXY -NO_WMAP -LARGEQUERY -NEG.SCORES=0 -WAIT -LONGLOC -DEV.TIMEOUT=120
-WARN.TIMEOUT=30 -THREDS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :
1: gb.ba:*
2: gb.htg:*
3: gb.in:*
4: gb.om:*
5: gb.ov:*
6: gb.pat:*
7: gb.ph:*
8: gb.pl:*
9: gb.pr:*
10: gb.ro:*
11: gb.sts:*
12: gb.sy:*
13: gb.un:*
14: gb.vl:*
15: em.ba:*
16: em.fun:*
17: em.hum:*
18: em.in:*
19: em.mu:*
20: em.om:*
21: em.or:*
22: em.ov:*
23: em.pat:*
24: em.ph:*
25: em.pl:*
26: em.ro:*
27: em.sts:*
28: em.un:*

29: em.vi:*
30: em.htg_hum:*
31: em.htg_inv:*
32: em.htg_other:*
33: em.htg_mus:*
34: em.htg_pin:*
35: em.htg_rod:*
36: em.htg_mam:*
37: em.htg_vrt:*
38: em.sy:*
39: em.htgo_hum:*
40: em.htgo_mus:*
41: em.htgo_other:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	193	100.0	169620	2 AC012674	AC012674 Homo sapi
C 2	193	100.0	199289	2 AC012378	AC012378 Homo sapi
C 3	129	66.8	152313	9 AL591594	AL591594 Human DNA
C 4	129	66.8	163314	2 AL590557	AL590557 Homo sapi
C 5	127	65.8	143372	9 AL137847	AL137847 Human DNA
C 6	117	60.6	148290	9 AC107979	AC107979 Homo sapi
C 7	117	60.6	165649	9 AC103996	AC103996 Homo sapi
C 8	117	60.6	169861	2 AC126324	AC126324 Homo sapi
C 9	117	60.6	192826	9 AC090762	AC090762 Homo sapi
C 10	116	60.1	131347	2 AC002421	AC002421 Homo sapi
C 11	116	60.1	170623	2 AL391375	AL391375 Human DNA
C 12	114	59.1	68314	2 AC126345	AC126345 Homo sapi
C 13	114	59.1	143717	2 AC069245	AC069245 Homo sapi
C 14	114	59.1	152354	2 AC016472	AC016472 Homo sapi
C 15	114	59.1	160066	2 AC027295	AC027295 Homo sapi
C 16	114	59.1	163475	9 AC078815	AC078815 Homo sapi
C 17	114	59.1	200543	2 AC016726	AC016726 Homo sapi
C 18	113	58.5	57662	2 AC107969	AC107969 Homo sapi
C 19	113	58.5	98360	9 HSD1247C2	AL049713 Human DNA
C 20	113	58.5	145264	9 AC107939	AC107939 Homo sapi
C 21	112	58.0	147820	9 AC092837	AC092837 Homo sapi
C 22	112	58.0	199038	2 AC116565	AC116565 Homo sapi
C 23	109	56.5	86314	9 AC109592	AC109592 Homo sapi
C 24	109	56.5	159475	2 AC021378	AC021378 Homo sapi
C 25	109	56.5	176689	9 AL162414	AL162414 Human DNA
C 26	108	56.0	134760	9 AC099484	AC099484 Homo sapi
C 27	108	56.0	166706	9 AC068875	AC068875 Homo sapi
C 28	108	56.0	207408	2 AC068618	AC068618 Homo sapi
C 29	108	56.0	207548	9 AC087283	AC087283 Homo sapi
C 30	107	55.4	103138	2 AC115094	AC115094 Homo sapi
C 31	107	55.4	143200	9 AC008413	AC008413 Homo sapi
C 32	107	55.4	231948	2 AC113346	AC113346 Homo sapi
C 33	106.5	55.2	146059	2 AC019030	AC019030 Homo sapi
C 34	106	54.9	177447	2 AC104687	AC104687 Homo sapi
C 35	106	54.9	236281	9 AC004673	AC004673 Homo sapi
C 36	105	54.4	80417	9 AL512659	AL512659 Human DNA
C 37	105	54.4	108248	9 HS326T13	AL022158 Homo sapi
C 38	105	54.4	162778	9 AC009559	AC009559 Homo sapi
C 39	104	53.9	24725	9 AL392090	AL392090 Human DNA
C 40	104	53.9	120418	2 AP002509	AP002509 Homo sapi
C 41	104	53.9	147288	9 AL355812	AL355812 Human DNA
C 42	104	53.9	163231	9 AL162575	AL162575 Human DNA
C 43	104	53.9	175832	9 AC023080	AC023080 Homo sapi
C 44	104	53.9	187847	9 AC023155	AC023155 Homo sapi
C 45	102	52.8	66489	2 AC100766	AC100766 Homo sapi

RESULT 1

ALIGNMENTS

AC012674/c
LOCUS AC012674 169620 bp DNA linear HTG 07-SEP-2000
DEFINITION Homo sapiens chromosome 3 clone RP1-458H3, WORKING DRAFT SEQUENCE.
AC012674
18 unordered pieces.
AC012674.10 GI:9719580
VERSION HTG: HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS Homo sapiens.
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Munnly,D.M., Adams,C., Bailey,M., Barbara,J., Blankenburg,K.,
1 (bases 1 to 169620)
Bodell,B., Bouck,J., Bowie,S., Brooks,A., Bunay,C., Bunac,C.,
Burkett,C., Burrows,J., Carter,M., Chacko,J., Chen,Z., Cox,C.,
David,R., Delgado,O., Deshazo,D., Ding,Y., Domah-Rashid,N.,
Dugan-Rocha,S., Durbin,K.J., Fernandez,C., Ferraigut,D.,
Forcum-Tansey,J., Franz,P., Ganesha,R., Gorrell,J.H., Gorrell,L.L.,
Guevara,W., Hosak,H., Jackson,L.E., Hodgson,A., Hogues,M.,
Holloway,C., Harris,K., Hernandez,J., Jackson,L., Jia,Y., Jones,M.,
Kelly,S., Kondejewski,N., Kong,Y., Kovar,C., Leal,B., Li,Z.,
Lichter,O., Liu,J., Liu,W., Logan,O., Lozano,R.J., Lu,J.,
Lucier,R., Martin,R., Martinez,C., McLeod,M.P., Mei,G., Morgan,M.,
Morris,S., Nash,S., Nelson,A., Nguyen,R., Nguyen,N., Nguyen,S.,
Oswall,G., Parish,B., Paxton,S., Payton,B., Perez,L., Pu,L.L.,
Quiles,M., Reiter,D., Rives,M., Samuel,S., Say,J., Scherer,S.,
Shah,E., Shen,H., Simon,M., Sparks,A., Stamps,A., Sugeng,S.,
Tabors,P., Taylor,T., Vasquez,L., Vinson,R., Vo,O., Wahab,M.,
Watlington,S., Weinstein,G., Weinstein,I.R., Williamson,A.,
Worley,K., Wren,J., Wrenford,G., Yu,W., Zhou,X., Nelson,D., and
Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley,K.C.
Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HMOG
Center clone name: RP1-458H3
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-contris estimation
Estimated insert size: 171608; agarose-1p estimation
Quality coverage: 3.9x in Q20 bases; agarose-1p estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contris estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank-draft-data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1 28689: contig of 28689 bp in length
* 28690 28789: gap of unknown length
* 50832 50832: contig of 22043 bp in length
* 50833 50932: gap of unknown length
* 50933 69144: contig of 18212 bp in length

* 69145 69244: gap of unknown length
* 69245 84204: contig of 14660 bp in length
* 84205 84304: gap of unknown length
* 84305 94667: contig of 10463 bp in length
* 94668 94767: gap of unknown length
* 94768 107261: contig of 12494 bp in length
* 107262 107361: gap of unknown length
* 107362 117550: contig of 10189 bp in length
* 117551 117650: gap of unknown length
* 117651 126939: contig of 9289 bp in length
* 126940 127039: gap of unknown length
* 127040 135040: contig of 8001 bp in length
* 135041 135140: gap of unknown length
* 135141 141639: contig of 6499 bp in length
* 141640 141739: gap of unknown length
* 141740 149558: contig of 7819 bp in length
* 149559 149658: gap of unknown length
* 149659 154562: contig of 4904 bp in length
* 154563 154662: gap of unknown length
* 154663 158987: contig of 4325 bp in length
* 158988 159087: gap of unknown length
* 159088 162376: contig of 3289 bp in length
* 162377 162476: gap of unknown length
* 162477 165191: contig of 2715 bp in length
* 165192 165291: gap of unknown length
* 165292 167173: contig of 1882 bp in length
* 167174 167273: gap of unknown length
* 167274 168393: contig of 1120 bp in length
* 168394 168493: gap of unknown length
* 168494 169620: contig of 1127 bp in length.
Location/Qualifiers
1. 169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP1-458H3"

BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others
ORIGIN

Alignment Scores:
Prid. No.: 9,18e-18 Length: 169620
Score: 193.00 Matches: 37
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 2 Gaps: 0

US-09-513-999c-copy_1_37 (1-37) x AC012674 (1-169620)

QY 1 MetGLYGISeRPhEaLAlEaUGInAaSPSeRPhSeSeRLeUGInGLyLeuLeUGLyPro 20
|||||
Db 87441 ATGGCGGAGTCTTTTCCTTCGAGGATTTCTTTTATCTTTTCGAGGACTTCTGGGGCCG 87382
|||||
QY 21 GLuTYrValLySLeUleUGLyLeUCySValCySLeUSerGLyCySerThr 37
|||||
Db 87381 GAGTATGTAAGACTCTGGGTCCTGTGTGTGTCCTGAGGCTGCTACT 87331
|||||

RESULT 2

AC012378 199289 bp DNA linear PRI 09-AUG-2001
LOCUS AC012378 Homo sapiens chromosome 15 clone RP11-420M1 map 15q21.3, complete
DEFINITION sequence.
AC012378
AC012378
VERSION AC012378.10 GI:15145648
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS 1 (bases 1 to 199289)
Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
Pate,D., and Hood,L.

TITLE Sequencing of human chromosome 15 D15S146-D15S117 region
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 199289)
AUTHORS Rowen L., Madan A., Qin S., Abbasi N., Baradarani L., Birditt B., Bloom S., Dors M., Dickhoff R., Harrison G., James R., Madan A., Owen M.P., Ratcliffe A., Shafer T. and Hood L.
JOURNAL Direct Submission
TITLE Submitted (26-OCT-1999) Multimegabase Sequencing Center, University of Washington, PO Box 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 199289)
AUTHORS Rowen L., Madan A., Qin S., Baradarani L., Birditt B., Bloom S., Burke J., Dors M., Fleetwood P., Kaur A., Madan A., Nesbitt R., Pate D. and Hood L.
JOURNAL Direct Submission
TITLE Submitted (09-AUG-2001) Multimegabase Sequencing Center, Institute for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA 98103, USA
COMMENT On Aug 9, 2001 this sequence version replaced gi:13775292.
----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leetowensystemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399

FEATURES
source
Note: Data from overlapping clones AC022083 [Drafting center: UWMSC], AC012674 [Drafting center: BCM] and AC009997 [drafting center: UWMSC] were added for finishing.
1. 199289
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15q21.3"
/clone="RP11-420M1"
/clone_lib="RPCT human bac library 11"
/note="This clone overlaps CTD-2137J4 AC022083, RP11-458H3 AC012674 and RP11-291H24 AC009997. Data from overlapping BACs were added and the consensus sequence determined from RP11-420M1 to the extent possible."
1. 165029
/note="Overlap with RP11-458H3, AC012674"
1. 11647
/note="Overlap with CTD-2137J4, AC022083"
49933. 50105
/note="Low quality data"
52173. 52233
/note="Low quality data"
52381. 52385
/note="Low quality data"
52412. 52483
/note="Low quality data"
52541. 52570
/note="Low quality data"
53242. 53257
/note="Low quality data"
53940. 54690
/note="Sequence data generated from subcloned PCR product"
72103. 72105
/note="Low quality data"
1503032. 155038
/note="Low quality data"
166787. 199289
/note="Overlap with RP11-291H24, AC009997"
BASE COUNT 63165 a 40338 c 36951 g 58835 t
ORIGIN
Alignment Scores:

Pred. No.: 1.08e-17 Length: 199289
Score: 193.00 Matches: 37
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0
US-09-513-999c-copy_1_37 (1-37) x AC012378 (1-199289)
QY 1 Merc1yGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyPro 20
Db 14756 ATGGTGGAATCTTTGGCTTCAGGATCTTTTCATCTTGGAGGACTCTGGGCGG 14697
QY 21 G1utTyValIysLeuLeuGlyLeuCySValCySLeuSerGlyCySerThr 37
Db 14696 GAGTATGTAACCTCGGCTCTGTGTGTGCTAGTGGCTGCTCTACT 14646
RESULT 3
AL591594/c 152313 bp DNA linear PRI 16-NOV-2001
LOCUS Human DNA sequence from clone RP11-424N15 on chromosome 1, complete
DEFINITION
ACCESSION AL591594 GI:16944148
VERSION AL591594.9
KEYWORDS HTG.
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 152313)
AUTHORS Almeida J.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humgeny@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk
On Nov 15, 2001 this sequence version replaced gi:1605741.
During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TrEMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C-elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr1
RP11-424N15 is from the library RPCT-11.2 constructed by the group of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACe3.6
This sequence is the entire insert of clone RP11-424N15 The true right end of clone RP11-518J10 is at 64264 in this sequence.
FEATURES
source
1. 152313
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-424N15"
/clone_lib="RPCT-11.2"
/note="Sequence from overlapping clone RP11-24C8
18821. 18877

```

BASE COUNT      45419 a 29865 c 29040 g 47988 t
ORIGIN
(ALT590557). Assembly confirmed by restriction digest."

Alignment Scores:
Pred. No.:          5,03e-08           Length:          152313
Score:              129.00             Matches:            23
Percent Similarity: 92.86%             Conservative:       3
Best local Similarity: 82.14%           Mismatches:         2
Query Match:        66.84%             Indels:             0
DB:                 9                  Gaps:               0

US-09-513-999C-7869_COPY_1_37 (1-37) x ALT591594 (1-152313)
OY      9   AspSerPheSerLeuIleuGlyLeuLeuAlpProGlyValIlysLeuIleuGlyLeu 28
||| |||||:::||: |||||:::||:|||||:::||:|||||:::||:|||||:::||:|||||
Db 100967  GACCAATTTTCACCTTGCGACGGGTCTTGGTCCACAGTAATAAATCTCGGACCCTC 100908

OY      29  CysValCysLeuSerGlyCysSer 36
|||||TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT
|||00907 TCGCTGTGCCTGAGTGGCTGCTCT 100884

RESULT 4
AL590557 LOCUS
DEFINITION Homo sapiens chromosome 1 clone RP11-24C8, *** SEQUENCING IN PROGRESS ***, 8 unordered pieces.
AL590557 AL590557 8 GI:13992136
HTG: HTGS_PHASE1, HTGS_CANCELLED.
human.
Organism Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 163314)
McLay, K.
Direct Submission
Submitted (20-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
On May 8, 2001 this sequence version replaced gi:13990622.
----- Genome Center -----
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
----- Project Information -----
Center project name: ba24c8
----- Summary Statistics -----
Sequencing program: XGAP4: version 4.5
Sequencing vector: plasmid: L08752; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 161385 bases at least Q40
Consensus quality: 161940 bases at least Q30
Consensus quality: 162264 bases at least Q20
Insert size: 162614; sum-of-contigs
Insert size: 164357; 4.9% error; agarose-fp
Quality coverage: 6.63x in Q20 bases; sum-of-contigs quality
coverage: 6.67x in Q20 bases; agarose-fp
----- NOTE: This is a 'working draft' sequence. It currently consists of 8 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as * runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
*****
1 16091: contig of 16091 bp in length
* 16092 16191: gap of 100 bp
* 16192 51184: contig of 34993 bp in length
* 51185 51284: gap of 100 bp
* 51285 67941: contig of 16657 bp in length

```

```

*      67942 68041: gap of 100 bp
*      68042 103921: contig of 35880 bp in length
*      103922 104021: gap of 100 bp
*      104022 107020: contig of 2999 bp in length
*      107021 107120: gap of 100 bp
*      107121 123748: contig of 16628 bp in length
*      123749 123848: gap of 100 bp
*      123849 137091: contig of 13243 bp in length
*      137092 137191: gap of 100 bp
*      137192 163314: contig of 26123 bp in length.
FEATURES
      source
          1..163314
              /organism="Homo sapiens"
              /db_xref="taxon:9606"
              /chromosome="1"
              /clone="RP11-24C8"
              /clone.lib="RPC1-11.1"
          1..16091
              /note="assembly:fragment:00603"
              /fragment_chain:1"
          16192..51184
              /note="assembly:fragment:01328"
              /fragment_chain:1"
          51285..67941
              /note="assembly:fragment:01579"
              /fragment_chain:1"
          68042..103921
              /note="assembly:fragment:00704"
              /fragment_chain:1"
          104022..107020
              /note="assembly:fragment:02225"
              /fragment_chain:1"
          107121..123748
              /note="assembly:fragment:02172"
              /fragment_chain:1"
          123849..137091
              /note="assembly:fragment:01998"
              /fragment_chain:1"
          137192..163314
              /note="assembly:fragment:03121"
              /note="assembly:fragment:03121"
BASE COUNT  51215 a 31325 c 32024 g 48050 t 700 others
ORIGIN
Alignment Scores:
Pred. No.:      5,39e-08      Length:      163314
Score:          129..00      Matches:      23
Percent Similarity: 92.86%      Conservative: 3
Best Local Similarity: 82.14%      Mismatches:  2
Query Match:      66.84%      Indels:      0
DB:              2      Gaps:      0
US-09-513-999C-7869_COPY_1_37 (1-37) x AL590557 (1-163314)
QY      9  AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTyrValLysLeuLeuGlyLeu 28
      |||
      |||
      |||
Db  55961  GACAAATTATTTTCCATTTGGCAGGGGTCCGCGGTCCACAGTATGTAACCTCCTGGGCCCTC 56020
      |||
      |||
      |||
QY      29  CysValGlySerLeuSerGlyCysSer 36
      |||
      |||
      |||
Db  56021  TGGCTGTGCTGTAGTGCTGCTCT 56044
      |||
      |||
      |||
RESULT 5
LOCUS      AL137847/c 143372 bp DNA linear PRI 16-NOV-2001
DEFINITION Human DNA sequence from clone RP11-439K3 on chromosome 9q22.2-31.1.1.
complete sequence.
ACCESSION  AL137847
VERSION    AL137847.12 GI:16973786
KEYWORDS   HTG.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

```


ACCESSION	AC103996	GI:21637504
VERSION	AC103996.7	
KEYWORDS	HTG.	
SOURCE	human.	
ORGANISM	Homo sapiens	

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE	AUTHORS	TITLE
1 (bases 1 to 165649)	Birren, B., Nusbaum, C. and Lander, E.	Homo sapiens chromosome 15, clone RP11-76E17

UNPUBLISHED
2 (bases 1 to 165649)
REFERENCE
AUTHORS

Barren,B., Linton,L., Nisbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barra,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepeil,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Deatellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heatford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., Lacroque,K.,
Lamatares,R., Landers,T., Lehoczyzy,T., Levine,R., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Margus,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., McSheeters,R., Meldrum,J.,
Meneses,L., Mihoiva,T., Mienga,P., Murphy,T., Naylor,J., Nguyen,C.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Rella,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Roselli,M., Roy,A., Santos,R., Schauer,S., Schupbach,R.,
Seaman,S., Severy,P., Spencer,B., Strange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zambek,L., Zlimer,A. and Zody,M.

JOURNAL
 submitted (01-DEC-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 165649)
 REFERENCE

TITLE Direct Submission
JOURNAL Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE 4 (bases 1 to 165649)

Allen, B., Nussbaum, C., Langer, E., Allia, Allen, N., Anderson, S., Barnes, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalil, B., Camarata, J., Chang, J., Chazaro, B., Choepel, Y., Collimore, A., Cook, A., Cooke, P., DeArillano, K., Dewar, K., Diaz, J. S., Dodge, S., Fero, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamel, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, C., Maclean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Melgrim, J., Meunier, L., Minova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicot, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Punniah, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,

TITLE
 JOURNAL
 COMMENT

Roman, J., Roy, A., Schauer, S., Schuppback, R., Seaman, S., Severy, P.,
 Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas,
 Tefayeh, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zalnoun, J.,
 Zembek, L., Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genomem
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Jul 1, 2002 this sequence version replaced g1:2159291.

FEATURES	
Source	Location/Qualifiers
	1. 165649
	/organism="Homo sapiens"
	/db_xref="taxon:9606"
	/chromosome="15"
	/map="15"
	/clone="RP11-76E17"
	/clone.lib="RPC1-11
repeat_region	complement(2..865)
	/rpt_family="L1PA13"
repeat_region	complement(864..894)
	/rpt_family="L1PA13"
repeat_region	complement(895..1248)
	/rpt_family="HHE1A"
repeat_region	complement(1249..1504)
	/rpt_family="L1PA13"
repeat_region	1505..1626
	/rpt_family="AluSx"
repeat_region	1627..1659
	/rpt_family="CAAA)n"
repeat_region	1660..1831
	/rpt_family="AluSx"
repeat_region	complement(1832..3132)
	/rpt_family="L1PA13"
repeat_region	complement(3140..3699)
	/rpt_family="L1MC"
repeat_region	complement(4461..4755)
	/rpt_family="AluSx"
repeat_region	5027..5397
	/rpt_family="L2"
repeat_region	complement(5650..6330)
	/rpt_family="L1ME1"
repeat_region	complement(6343..6460)
	/rpt_family="L1ME1"
repeat_region	7214..7319
	/rpt_family="L1MC3"
repeat_region	7354..7655
	/rpt_family="L1MC3"
repeat_region	7659..7743
	/rpt_family="L1PA10"
repeat_region	complement(7745..7843)
	/rpt_family="AluSv4"
repeat_region	7844..14156
	/rpt_family="L1PA10"
repeat_region	14198..14328
	/rpt_family="AluY"
repeat_region	14340..14503
	/rpt_family="(TA)n"
repeat_region	14507..15032
	/rpt_family="L1MC3"
repeat_region	15036..15220
	/rpt_family="L1MC3"

```

repeat_region complement(15218..16449)
repeat_region /rpt_family="L1PA4"
16450..18154
repeat_region /rpt_family="L1PA4"
18155..19210
repeat_region /rpt_family="L1MC3"
19211..19266
repeat_region /rpt_family="(TA)n"
19267..19372
repeat_region /rpt_family="L1MC3"
19401..19476
repeat_region /rpt_family="(TTATA)n"
19483..19546
repeat_region /rpt_family="(CATATA)n"
19596..19625
repeat_region /rpt_family="AT-rich"
19696..19750
repeat_region /rpt_family="GA-rich"
19752..19920
repeat_region /rpt_family="L1MD3"
19994..20102
repeat_region /rpt_family="L2"
20049..20114
unsure /note="single clone coverage"
repeat_region /rpt_family="MIR3"
20485..20655
repeat_region complement(21285..21441)
/rpt_family="MIR"
21496..21717
repeat_region /rpt_family="L2"
/rpt_family="MIR"
21720..21796
repeat_region /rpt_family="MIR"
complement(21943..22131)
/rpt_family="MIR"
repeat_region 23082..23195
/rpt_family="L2"
23198..23248
repeat_region /rpt_family="GA-rich"
23267..23333
repeat_region /rpt_family="(CAT)n"
complement(24465..24833)
/rpt_family="M1T1A2"
26142..26334
repeat_region /rpt_family="MIR"
/rpt_family="MIR"
complement(27659..27811)
repeat_region /rpt_family="MIR"
complement(28811..28880)
repeat_region /rpt_family="MIR"
complement(29026..29269)
/rpt_family="MIR"
complement(29673..29706)
/rpt_family="MSTB"

Alignment Scores:
Pred. No.: 3.74e-06 Length: 165649
Score: 117.00 Matches: 21
Percent Similarity: 85.71% Conservative: 4
Best Local Similarity: 75.01% Mismatches: 4
Query Match: 60.62% Indels: 0
DB: 9 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC103996 (1-165649)
OY 9 AspserPheserSerLeuGlnGlyLeuLeuGlyProGluTyrValValysLeuLeuGlyLeu 28
DB 8153 GATGATATCTTCACCTCACTGCGGATCTCTGGGCGACAGATATATAAACTCTCGGCTCC 8094
OY 29 CysValCysLeuSerGlyCysSer 36
DB 8093 TGTGTGTGCTGCTGAGTGGCTGCTCA 8070
RESULT 8
AC126324/c

```

```

LOCUS AC126324 169861 bp DNA linear HTG 06-AUG-2002
DEFINITION Homo sapiens chromosome 11 clone RP11-358H20 map 11. *** SEQUENCING
IN PROGRESS ***; 2 ordered pieces.
ACCESSION AC126324
VERSION AC126324.2 GI:22123018
KEYWORDS HTG; HTGS_PHASE2; HTGS_FULFILLTOP; HTGS_ACTIVEFIN.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 169861)
Birren,B., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 11, clone RP11-358H20
Unpublished
2 (bases 1 to 169861)
Birren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Bookhalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,D., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schnpack,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Willson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (05-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 169861)
Birren,B., Bastien,V., Bloom,T., Boguslavsky,L., Bookhalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,D., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schnpack,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Willson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (06-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 6, 2002 this sequence version replaced g1:21699255.
All repeats were identified using RepeatMasker:
Smit,A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RN/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIRB
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L27749
Center clone name: 358_H_20

```

* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. Gaps between the contigs

repeat_region	7329. .7446	/rpt_family="MIR"	complement(8147. .8452)
repeat_region		/rpt_family="AluIo"	8540. .8639
repeat_region		/rpt_family="MER45"	complement(8832. .8916)
repeat_region		/rpt_family="L2"	10391. .10599
repeat_region		/rpt_family="MER3"	11688. .11828
repeat_region		/rpt_family="L1MC/D"	12029. .12078
repeat_region		/rpt_family="AT_rich"	12092. .12447
repeat_region		/rpt_family="MHELIC"	complement(13616. .13751)
repeat_region		/rpt_family="MIR3"	complement(13358. .14142)
repeat_region		/rpt_family="MIR"	complement(155277. .15553)
repeat_region		/rpt_family="MER8"	complement(15511. .16005)
repeat_region		/rpt_family="MIR"	complement(16101. .16440)
repeat_region		/rpt_family="L3"	16920. .16958
repeat_region		/rpt_family="TCCC0"	complement(17145. .17444)
repeat_region		/rpt_family="AluSx"	complement(18418. .19953)
repeat_region		/rpt_family="L1MC"	complement(19978. .20262)
repeat_region		/rpt_family="L1MC"	complement(20788. .20794)
repeat_region		/rpt_family="L1MC"	complement(2082. .21097)
repeat_region		/rpt_family="L1MC"	complement(21445. .21743)
repeat_region		/rpt_family="L1MC"	22599. .23518
repeat_region		/rpt_family="L1MC"	23527. .23901
repeat_region		/rpt_family="L1MC"	complement(23237. .24026)
repeat_region		/rpt_family="MSTB1"	complement(24027. .24256)
repeat_region		/rpt_family="MER30"	complement(24257. .24557)
repeat_region		/rpt_family="MSTB1"	24563. .24594
repeat_region		/rpt_family="AT_rich"	complement(25258. .25639)
repeat_region		/rpt_family="L1MC"	complement(25688. .25838)
repeat_region		/rpt_family="L1MC"	26167. .26506
repeat_region		/rpt_family="THE1B"	26746. .27094
repeat_region		/rpt_family="Tiger2a"	27095. .27184
repeat_region		/rpt_family="MAD1"	27185. .27279
repeat_region		/rpt_family="Tiger2a"	28321. .28363
repeat_region		/rpt_family="(TG)n"	29172. .29333
repeat_region		/rpt_family="MIR"	30664. .30898
repeat_region		/rpt_family="L1MB8"	31577. .31598
repeat_region		/rpt_family="AT_rich"	32378. .32475

```
repeat_region /rpt_family="Cr-rich" 34655.34960
repeat_region /rpt_family="AluY" complement(33296.35589)
repeat_region /rpt_family="MER5B" complement(35419.35517)
repeat_region /rpt_family="L1MC4" complement(33518.35823)
repeat_region /rpt_family="AluSX" complement(33824.36030)
repeat_region /rpt_family="L1MC4" 36682.36758
repeat_region /rpt_family="MER5A" 36975.37160
repeat_region /rpt_family="MER5A" /rpt_family="MER5A" 38140.38353
repeat_region /rpt_family="MER96B" 38633.38656
repeat_region /rpt_family="TTTA1n" complement(38657.40056)
repeat_region /rpt_family="L1PAs" complement(40060.40250)
```

Alignment Scores:	
Pred. No.:	4 34e-06
Score:	117.00
Percent Similarity:	85.71%
Best Local Similarity:	75.00%
Query Match:	60.62%
DB:	9
	Gaps: 0
	Mismatches: 4
	Conservative: 3
	Matches: 21
	Indels: 0
	Gaps: 0
	Length: 192826

US-09-513-999C-7869_COPY_1_37 (1-37) x AC090762 (1-192826)

QY	9	AsSerPheSerLeuGlnGlyLeuLeuGlnIprGlnIyrValLysLeuLeuGlyLeu	28
Db	190989	GATGATCTTTCACCTCAGCGGGATCTCTGGGCCACAGTATATATAAACTCTGGGCTC	190930
QY	29	CysValCysLeuSerGlyCysSer	36
Db	190929	TGTGTGCTCTGAGTGGCTCTCA	190906

RESULT 10	
AC002421	
LOCUS	131347 bp DNA linear HTG 13-JUN-2002
DEFINITION	Homo sapiens chromosome X clone pMXD1, ** SEQUENCING IN PROGRESS
	*** 4 unordered pieces.

ACCESSION	AC002421
VERSION	AC002421.2
KEYWORDS	GI:21405641
	HTG; HTGS_PHASE1.

ORGANISM Homo sapiens

REFERENCE
1 (bases 1 to 131347)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

AUTHORS
Chen, E., Brownstein, B.H., States, D.J., Schlessinger, D. and
Mazzarella, R.

TITLE	Direct Submission
JOURNAL	Unpublished (1997)

REFERENCE
AUTHORS 2 (bases 1 to 131347)
Brownstein, B.H., States, D.J. and Mazzarella, R.

TITLE Direct Submission
JOURNAL Submitted (12-AUG-1997) Center for Genetics in Medicine, Box 8232

Washington University School of Medicine, 4566 Scott Avenue, St. Louis, MO 63110, USA
On Jun 13, 2002 this sequence version replaced gi:2323248.

Current status of this project is available at
'http://genome.wustl.edu/cgm/seq_projects.html
Submitted by:

Elison Chen,
Advanced Center for Genetic Technology,
Applied Biosystems Division of Perlin Elmer Corp.,
850 Lincoln Center Drive,
Foster City, CA 94404 USA

e-mail: ellson@genseq.apldbio.com

and

Buddy Brownstein,

Center for Genetics in Medicine,
Washington University School of Medicine, Box 8232
4566 Scott Avenue,
St. Louis, MO 63110, USA

e-mail: buddy@genetics.wustl.edu

and

David J. States,
Institute for Biomedical Computing
Washington University in St. Louis
700 South Euclid Ave.
St. Louis, MO 63108 USA

e-mail: states@dbc.wustl.edu.

* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 22595: contig of 22595 bp in length
* 22596 22695: gap of 100 bp
* 22696 66983: contig of 44288 bp in length
* 66984 67083: gap of 100 bp
* 67084 122847: contig of 55764 bp in length
* 122848 122947: gap of 100 bp
* 122948 131347: contig of 8400 bp in length.

FEATURES

source

1. 131347
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="PMXD1"

BASE COUNT 40590 a 23040 c 23923 g 43494 t 300 others

ORIGIN

Alignment Scores:

Pred. No.: 4.22e-06 Length: 131347
Score: 116.00 Matches: 21
Percent Similarity: 86.21% Conservative: 4
Best Local Similarity: 72.41% Mismatches: 4
Query Match: 60.10% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC002421 (1-131347)

OY 9 AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGlnTyrValIysLeuGlnGlyLeu 28

Db 103911 GATGGATCGCCACCTTCGTGGGAATCTGGGGCCAGAGTAAACTCTCGGCTCTC 103970

OY 29 CysValCysLeuSerGlyCysSerThr 37

Db 103971 TGTGTGTGCTGAGCAGCTGCTCTCT 103997

RESULT 11

AL391375

LOCUS AL391375 170623 bp DNA linear PRI 02-FEB-2001
DEFINITION Human DNA sequence from clone RP11-375A20 on chromosome X, complete
sequence.

ACCESSION AL391375 GI:12657182

VERSION

KEYWORDS

HTG

SOURCE

ORGANISM

Homo sapiens
human.
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE

AUTHORS

Chapman, J.

Direct Submission

JOURNAL

Submitted (02-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone

COMMENT

requests: clonerequest@sanger.ac.uk

On Feb 2, 2001 this sequence version replaced gi:11414631.

During sequence assembly data is compared from overlapping clones,
where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.

This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated repeat sequence elements. Where the sequence is
ambiguous, there is an annotation using the 'unsure' feature key.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/projects/C.elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/ChrX

RP11-375A20 is from the library RPCI-11.2 constructed by the group
of Pletier de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6

FEATURES

source

1. 170623
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="RP11-375A20"
/clone_11b="RPCI-11.2"
71147..71285
misc_feature
/note="Single clone region. Assembly confirmed by
restriction digest data."

BASE COUNT 53145 a 29784 c 30919 g 56775 t

ORIGIN

Alignment Scores:

Pred. No.: 5.47e-06 Length: 170623
Score: 116.00 Matches: 21
Percent Similarity: 86.21% Conservative: 4
Best Local Similarity: 72.41% Mismatches: 4
Query Match: 60.10% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AL391375 (1-170623)

OY 9 AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGlnTyrValIysLeuGlnGlyLeu 28

Db 99926 GATGGATCGCCACCTTCGTGGGAATCTGGGGCCAGAGTAAACTCTCGGCTCTC 99985

OY 29 CysValCysLeuSerGlyCysSerThr 37

Db 99986 TGTGTGTGCTGAGCAGCTGCTCTCT 100012

RESULT 12

AC126345/c

LOCUS AC126345 68314 bp DNA linear HTG 30-JUL-2002
DEFINITION Homo sapiens chromosome 11 clone RP11-100E23 map 11, LOW-PASS
SEQUENCE SAMPLING.

ACCESSION

AC126345 GI:22004319

VERSION

KEYWORDS

HTG

SOURCE

ORGANISM

Homo sapiens
human.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata: Euteleostomi;
Mammalia: Eutheria: Primates; Catarrhini: Hominoidea: Homo.
1 (bases 1 to 68314)
Birren,B., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 11, clone RP11-100E23
Unpublished
2 (bases 1 to 68314)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhalter,B.,
Camarata,J., Chang,U., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardina,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Hornton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Miengs,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Nordu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,B., Schuppback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Strange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (05-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 68314)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhalter,B.,
Camarata,J., Chang,U., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardina,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Hornton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Miengs,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Nordu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,B., Schuppback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Strange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (30-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 30, 2002 this sequence version replaced gi:21699290.
ALL repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

Project Information
Center project name: L27743
Center clone name: 100_E_23

* NOTE: This record contains 84 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will

* be preserved. 728: contig of 728 bp in length
* 1
* 729 828: gap of 100 bp
* 829 1544: contig of 716 bp in length
* 1545 1644: gap of 100 bp
* 1645 2360: contig of 716 bp in length
* 2361 2460: gap of 100 bp
* 2461 3152: contig of 692 bp in length
* 3153 3252: gap of 100 bp
* 3253 3960: contig of 708 bp in length
* 3961 4060: gap of 100 bp
* 4061 4774: contig of 714 bp in length
* 4775 4874: gap of 100 bp
* 4875 5583: contig of 709 bp in length
* 5584 5683: gap of 100 bp
* 5684 6389: contig of 706 bp in length
* 6390 6489: gap of 100 bp
* 6490 7213: contig of 724 bp in length
* 7214 7313: gap of 100 bp
* 7314 8019: contig of 706 bp in length
* 8020 8119: gap of 100 bp
* 8120 8851: contig of 732 bp in length
* 8852 8951: gap of 100 bp
* 8952 9622: contig of 671 bp in length
* 9623 9722: gap of 100 bp
* 9723 10434: contig of 712 bp in length
* 10435 10534: gap of 100 bp
* 10535 11257: contig of 723 bp in length
* 11258 11357: gap of 100 bp
* 11358 12070: contig of 713 bp in length
* 12071 12170: gap of 100 bp
* 12171 12895: contig of 725 bp in length
* 12896 12995: gap of 100 bp
* 12996 13714: contig of 719 bp in length
* 13715 13814: gap of 100 bp
* 13815 14524: contig of 710 bp in length
* 14525 14624: gap of 100 bp
* 14625 15335: contig of 711 bp in length
* 15336 15435: gap of 100 bp
* 15436 16165: contig of 730 bp in length
* 16166 16265: gap of 100 bp
* 16266 16986: contig of 721 bp in length
* 16987 17086: gap of 100 bp
* 17087 17817: contig of 731 bp in length
* 17818 17917: gap of 100 bp
* 17918 18633: contig of 716 bp in length
* 18634 18733: gap of 100 bp
* 18734 19459: contig of 726 bp in length
* 19460 19559: gap of 100 bp
* 19560 20266: contig of 707 bp in length
* 20267 20366: gap of 100 bp
* 20367 21090: contig of 724 bp in length
* 21091 21190: gap of 100 bp
* 21191 21901: contig of 711 bp in length
* 21902 22001: gap of 100 bp
* 22002 22721: contig of 720 bp in length
* 22722 22821: gap of 100 bp
* 22822 23530: contig of 709 bp in length
* 23531 23630: gap of 100 bp
* 23631 24336: contig of 706 bp in length
* 24337 24436: gap of 100 bp
* 24437 25167: contig of 731 bp in length
* 25168 25267: gap of 100 bp
* 25268 25989: contig of 722 bp in length
* 25990 26089: gap of 100 bp
* 26090 26816: contig of 727 bp in length
* 26817 26916: gap of 100 bp
* 26917 27618: contig of 702 bp in length
* 27619 27718: gap of 100 bp
* 27719 28439: contig of 721 bp in length
* 28440 28539: gap of 100 bp
* 28540 29257: contig of 718 bp in length
* 29258 29357: gap of 100 bp

```
* 29358 30053: contig of 696 bp in length
* 30054 30153: gap of 100 bp
* 30154 30856: contig of 703 bp in length
* 30857 30956: gap of 100 bp
* 30957 31680: contig of 724 bp in length
* 31681 31780: gap of 100 bp
* 31781 32509: contig of 729 bp in length
* 32510 32609: gap of 100 bp
* 32610 33331: contig of 722 bp in length
* 33332 33431: gap of 100 bp
* 33432 34153: contig of 722 bp in length
* 34154 34253: gap of 100 bp
* 34254 34964: contig of 711 bp in length
* 34965 35064: gap of 100 bp
* 35065 35759: contig of 695 bp in length
* 35760 35859: gap of 100 bp
* 35860 36566: contig of 707 bp in length
* 36567 36666: gap of 100 bp
* 36667 37385: contig of 719 bp in length
* 37386 37485: gap of 100 bp
* 37486 38202: contig of 717 bp in length
* 38203 38302: gap of 100 bp
* 38303 39019: contig of 717 bp in length
* 39020 39119: gap of 100 bp
* 39120 39833: contig of 714 bp in length
* 39834 39933: gap of 100 bp
* 39934 40657: contig of 724 bp in length
* 40658 40757: gap of 100 bp
* 40758 41471: contig of 714 bp in length
* 41472 41571: gap of 100 bp
* 41572 42284: contig of 713 bp in length
* 42285 42384: gap of 100 bp
* 42385 43100: contig of 716 bp in length
* 43101 43200: gap of 100 bp
* 43201 43926: contig of 726 bp in length
* 43927 44026: gap of 100 bp
* 44027 44745: contig of 719 bp in length
* 44746 44845: gap of 100 bp
* 44846 45552: contig of 707 bp in length
* 45553 45652: gap of 100 bp
* 45653 46336: contig of 684 bp in length
* 46337 46436: gap of 100 bp
* 46437 47128: contig of 692 bp in length
* 47129 47228: gap of 100 bp
* 47229 47951: contig of 723 bp in length
* 47952 48051: gap of 100 bp
```

Alignment Scores:
Query No.: 4.46e-06 Length: 68314
Percent Similarity: 114.00 Matches: 22
Best Local Similarity: 83.87% Conservative: 4
Query Match: 59.07% Mismatches: 5
Indels: 0
Gaps: 0

US-09-513-999c-7869_copy_1_37 (1-37) x AC126345 (1-68314)

QY 6 AAlaenGlaAspSerPheSerLeuGlnGlyLeuLeuGlyPProGluTyrValIysIeu 25

Db 43053 GCCCTGCGCTGCACTTGCACGACAGTTTGTTGGGATCCCTTGGCCGACGAGTTGTAAAGCTC 42994

QY 26 LeuGlyLeuCyValCysIeuSerGlyCysSer 36

Db 42993 CTGGGCTCTGTGTGTGTGTGACGACGCTCTCT 42961

RESULT 13

AC069245

LOCUS AC069245 143717 bp DNA linear HTG 09-MAY-2002

DEFINITION Homo sapiens chromosome 12 clone RP11-757N13, WORKING DRAFT

AC069245

AC069245.10 GI:20335692

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE Homo sapiens.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 143717)
AUTHORS Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C., Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Ayele,M., Banks,T., Barbarella,J., Benton,J., Bimaga,K., Blankenburg,K., Bonini,D., Bouck,J., Bowles,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C., Burck,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Z., Chowdhury,I., Christopoulos,C., Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Dublin,K.J., Earhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K., Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C., Hollins,B., Homs,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudan,S., Karlsson,E., Kelly,S., Khan,U., King,L., Kovach,J., Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L., Li,J., Li,Z., Lichtharge,O., Lieu,C., Liu,J., Liu,W., Lousegod,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E., Massey,E., Mawhney,E., McLeod,M.P., Meador,M., Mel,G., Metzger,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Nohabhat,K., Morgan,M., Morris,S., Nguyen,N., Nickerson,E., Nwokewho,S., Oguh,M., Okunnu,G., Orangunye,N., Oyiedo,R., Pace,A., Payton,B., Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojudoan,I., Rolfe,M., Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shooshari,N., Sisson,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,H., Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S., Usmani,K., Vasquez,L., Vera,Y., Villalón,D., Vinson,R., Wang,Q., Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S., Williams,G., Williamson,A., Wleczky,R., Wooden,S., Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D., Weinstein,G., and Gibbs,R.

TITLE Direct Submission

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 143717)

AUTHORS Worley,K.C.

JOURNAL Direct Submission

REFERENCE Submitted (22-MAY-2000)

AUTHORS Human Genome Sequencing Center, Department

JOURNAL of Molecular and Human Genetics, Baylor College of Medicine, One

COMMENT Baylor Plaza, Houston, TX 77030, USA

On Apr 28, 2002 this sequence version replaced gi:18449608.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: HBX

Center clone name: RP11-757N13

----- Summary Statistics

Sequencing vector: M13

Chemistry: Dye-primer Bodypy: 19% of reads

Chemistry: Dye-terminator Big Dye: 81% of reads

Assembly program: Phrap; version 0.990329


```
* 3710 3809: gap of 100 bp
* 3810 6730: contig of 2921 bp in length
* 6731 6830: gap of 100 bp
* 6831 9206: contig of 2376 bp in length
* 9207 9306: gap of 100 bp
* 9307 12163: contig of 2857 bp in length
* 12164 12265: gap of 100 bp
* 12264 15381: contig of 3118 bp in length
* 15382 15481: gap of 100 bp
* 15482 20100: contig of 4619 bp in length
* 20101 20200: gap of 100 bp
* 20201 23885: contig of 3685 bp in length
* 23886 23985: gap of 100 bp
* 23986 27226: contig of 3241 bp in length
* 27227 27326: gap of 100 bp
* 27327 32869: contig of 5543 bp in length
* 32870 32969: gap of 100 bp
* 32970 36903: contig of 3934 bp in length
* 36904 37003: gap of 100 bp
* 37004 44368: contig of 7366 bp in length
* 44370 44469: gap of 100 bp
* 44470 53277: contig of 8808 bp in length
* 53278 53377: gap of 100 bp
* 53378 71089: contig of 17712 bp in length
* 71090 71189: gap of 100 bp
* 71190 90983: contig of 19794 bp in length
* 90984 91083: gap of 100 bp
* 91084 112306: contig of 21223 bp in length
* 112307 112406: gap of 100 bp
* 112407 152354: contig of 39948 bp in length.
Location/Qualifiers
1. 152354
```

```
FEATURES
SOURCE
1. 152354
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-20p1"
/clone_id="RPC1-11 Human Male BAC"
```

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

```
BASE COUNT 42989 a 30395 c 30652 g 46711 t 1607 others
ORIGIN
```

```
Alignment Scores:
Pred. No.: 9.89e-06 Length: 152354
Score: 114.00 Matches: 20
Percent Similarity: 85.71% Conservative: 4
Best Local Similarity: 71.43% Mismatches: 4
Query Match: 59.07% Indels: 0
DB: 2 Gaps: 0
```

US-09-513-999c-7869_COPY_1_37 (1-37) x AC016472 (1-152354)

QY 9 AspSerPheSerSerLeuGlnGlyLeuGlnGlyProGlnItyrValIysLeuGlnGlyLeu 28

Db 117786 GATGATTTCCACCTGCTGCTGATCTCGTGGACACAGATATTAACATCATGCTCTC 117845

QY 29 CysValCysLeuSerGlyCysSer 36

Db 117846 TGTATGTGCTGACGACATGCTCT 117869

RESULT 15

AC027295

LOCUS

DEFINITION

AC027295

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 160066)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Ayalew,S.R., Banks,T.,
Barbarta,J., Benton,J., Bimsge,K., Blankenburg,K., Bonnin,D.,
Bouch,J., Bowie,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P.,
Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,
Carron,T.F., Carter,M., Cavazos,S.R., Checko,J., Chavez,D.,
Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C.,
Cleveland,C.D., Cox,C., Coyle,M.D., Dethorne,S.R., David,R.,
Dayila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,
DeLaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H.,
Douthwaite,K.J., Draper,H., Dugan-Hochs,S., Durbin,K.J.,
Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M.,
Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P.,
Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R.,
Gibrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K.,
Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J.,
Hernandez,O., Hodgson,A., Hogues,M., Holloway,C., Hollins,B.,
Honsli,F., Howard,S., Huber,J., Huijk,S., Hume,J., Jackson,L.E.,
Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S.,
Karlsom,E., Kelly,S., Khan,U., King,L., Korvah,J., Kovar,C.,
Kratovic,J., Kureshi,A., Landry,N., Deal,B., Lewis,L.C., Lewis,L.,
Li,J., Li,Z., Licharge,O., Lieu,C., Liu,J., Liu,W., Lounseged,B.,
Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J.,
Maheshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E.,
Massey,E., Mawhinney,E., McLeod,M.P., Meador,M., Mei,G., Metzger,M.,
Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S.,
Mosier,M., Neal,D., Newton,J., Newton,N., Nguyen,A., Nguyen,N.,
Nguyen,N., Nickerson,E., Nwokenwo,S., Ogih,M., Okwouu,G.,
Oragunye,N., Oviado,R., Pace,A., Payton,B., Peery,J., Perez,L.,
Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y.,
Rivers,M., Rojas,A., Rojubokan,I., Rolfe,M., Ruiz,S., Saverly,G.,
Scherer,S., Scott,G., Shen,H., Shoshchari,N., Sisson,I.,
Sodergren,E., Sonalke,T., Sparks,A., Stanley,R., Stone,H.,
Sutton,A., Syatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H.,
Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,
Usmanli,K., Vasquez,L., Vera,V., Villalobon,D., Vinson,R., Wang,Q.,
Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,
Williams,G., Williamson,A., Wleczky,R., Wooden,S., Worley,K.,

Wu.C., Wu.Y., Wu.Y.F., Zhou.J., Zorrilla.S., Nelson.D.,
Weinstock.G. and Gibbs.R.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 160066)
AUTHORS Worley.K.C.
TITLE Direct Submission
JOURNAL Submitted (30-MAR-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 160066)
Worley.K.C.
REFERENCE Direct Submission
TITLE Submitted (18-JUN-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jun 17, 2002 this sequence version replaced gi:20335550.

COMMENT

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Drafting Center Code: BCM

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: HATS

Center clone name: RP11-293C20

----- Summary Statistics

Sequencing vector: M13;

Chemistry: Dye-primer Bodipy: 42% of reads

Chemistry: Dye-terminator Big Dye: 58% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 144209 bases at least Q40

Consensus quality: 149815 bases at least Q30

Estimated insert size: 1574/3; sum-of-coverage estimation

Quality coverage: 4x in Q20 bases; sum-of-coverage estimation

* NOTE: Estimated insert size may differ from sequence length

* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 13 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 3659: contig of 3659 bp in length

* 3660 3759: gap of unknown length

* 3760 5914: contig of 2155 bp in length

* 5915 6014: gap of unknown length

* 6015 9826: contig of 3812 bp in length

* 9827 9926: gap of unknown length

* 9927 12721: contig of 2794 bp in length

* 12721 12820: gap of unknown length

* 12821 20586: contig of 7766 bp in length

* 20587 20686: gap of unknown length

* 20687 27322: contig of 6636 bp in length

* 27323 27422: gap of unknown length

* 27423 39529: contig of 12107 bp in length

* 39530 39629: gap of unknown length

* 39630 54210: contig of 14581 bp in length

* 54211 54310: gap of unknown length

* 54311 64148: contig of 9838 bp in length

* 64149 64248: gap of unknown length

* 64249 79424: contig of 15176 bp in length

* 79425 79524: gap of unknown length

* 79525 99643: contig of 20019 bp in length

* 99644 125582: gap of unknown length

* 125583 125682: contig of 25939 bp in length

* 125683 160066: gap of unknown length

* Location/Qualifiers

source

1..160066

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="12"

/clone="RP11-293C20"

BASE COUNT 47149 a 32911 c 32401 g 46353 t 1252 others

ORIGIN

Alignment Scores:

Pred. No.:

Score:

Percent Similarity:

Best Local Similarity:

Query Match:

DB:

US-09-513-999c-7869_copy_1_37 (1-37) x AC027295 (1-160066)

QY

9

AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTrpValLysLeuLeuGlyLeu

DB 49872

GATGATTTTCCACCTTCTGCGATCTCGGACACAGATATGTAACATCATGCTCTC

QY 29

CysValLysLeuSerGlyCysSer 36

DB 49932

TGTATGTGCTCCTGACGATGCTCT 49955

Search completed: April 24, 2003, 23:01:33

Job time : 1526 secs

GenCore version 5.1.4-p5_4578
Copyright (c) 1993 - 2003 Compugen Ltd.

SUMMARIES

OM protein - nucleic search, using frame_plus_p2n model

Run on: April 24, 2003, 22:31:13 ; Search time 152 Seconds
(without alignments)
548.184 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSGFALDSSFSIQGLGPEYKVLGLCYCLSGCST 37

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

atched: 2185239 segs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame+ p2n.model -DEV=xlh
-Q=/cgn2.1/USPTO/US09513999/rumat_18042003.170936_28347/app_query.fasta.1.199
-DB=N.Geneseq_101002 -QFMT=fastap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOPCTL=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOCCALIGN=200 -THR_SCORE=pcr -THR_MAX=100 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=pcr -NOR=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn.1.1.200.0@rnat.18042003.170936.28347 -NCPU=6 -ICPU=3
-NO_XLPPY -NO_WMAP -LARGQUEYRY -NEG_SCORES=0 -WAIT -LONGLOG -DEV_TIMEOUT=120
-NARN_TIMEOUT=30 -THRAADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :
N.Geneseq_101002:*
1: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1980.DAT:*
2: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1981.DAT:*
3: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1983.DAT:*
4: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1984.DAT:*
5: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1985.DAT:*
6: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1986.DAT:*
7: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1987.DAT:*
8: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1988.DAT:*
9: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1989.DAT:*
10: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1990.DAT:*
11: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1991.DAT:*
12: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1992.DAT:*
13: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1993.DAT:*
14: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1994.DAT:*
15: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1995.DAT:*
16: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1996.DAT:*
17: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1997.DAT:*
18: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1998.DAT:*
19: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1999.DAT:*
20: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2000.DAT:*
21: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001.DAT:*
22: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
23: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	DB ID	Description
1	193	100.0	447	21 AAC03794	Human secreted pro
2	193	100.0	447	21 AA242680	Human 5' EST isola
3	95	49.2	128600	24 ABR83461	Human CDNA differe
4	88.5	45.9	570	22 ABA63453	Human foetal liver
5	88.5	45.9	570	22 ABA30652	Probe #9118 for ge
6	88.5	45.9	570	22 AAK11985	Human brain expres
7	88.5	45.9	570	22 AAK37688	Human bone marrow
8	88.5	45.9	570	22 AAI18447	Probe #8380 for ge
9	88.5	45.9	570	22 AAI43563	Probe #12249 used
10	88.5	45.9	570	24 ABS11680	Human genome-deriv
11	88	45.6	1982	21 AAC68089	Human secreted pro
12	86	44.6	1299	24 AAS91827	DNA encoding novel
13	85	44.0	240	24 ABR25478	Human OREF polynuc
14	83	43.0	1909	23 AAS88434	DNA encoding novel
15	78	40.4	609	22 AAR98980	Human EST-derived
16	75	38.9	349	22 AAR65888	Novel Human polynu
17	71	36.8	11534	24 ABL32342	Human immune syste
18	69	35.8	2787	23 AAS68019	DNA encoding novel
19	69	35.8	2787	23 AAS69828	DNA encoding novel
20	67	34.7	1996	22 ABA18896	Human nervous syst
21	65	33.7	447	22 AAI16101	Human breast cance
22	65	33.7	447	22 AAI24945	Human breast cance
23	64	33.2	36901	20 AA23892	Murine LOBO genom
24	64	33.2	38886	20 AA23897	Murine LOBO genom
25	63	32.6	162	22 ABA75762	Human foetal liver
26	63	32.6	519	22 ABA63320	Human foetal liver
27	60.5	31.3	2730	22 ABA84293	Human EXCS encodin
28	60.5	31.3	24533	22 AAS27689	DNA encoding novel
29	59	30.6	481	23 ABV05461	Human prostate exp
30	59	30.6	560	22 AAK78945	Human immune/haema
31	58.5	30.3	27666	23 ABL08332	Drosophila melano
32	58	30.1	2085	20 AAX01362	Nucleobase permas
33	57.5	29.8	1605	24 ABL40748	Chicken hepatocarc
34	57	29.5	2743	22 AAK86888	Human immune/haema
35	57	29.5	7379	19 AA49653	Human SCL DNA. Ho
36	56.5	29.3	259	23 ABV05218	Human prostate exp
37	56.5	29.3	259	23 ABV14387	Human prostate exp
38	56.5	29.3	400	23 ABV35471	Human prostate exp
39	56.5	29.3	650	24 ABO56543	Human colon cancer
40	56.5	29.3	2076	23 AAS64519	DNA encoding novel
41	56	29.0	1226	22 AAK53271	Human polynucleoti
42	56	29.0	1380	24 ABR73394	Bacillus licheniflo
43	56	29.0	2015	22 AAR57417	Human stomach cell
44	56	29.0	2073	20 AAK87925	Human protease HPR
45	56	29.0	2118	22 AAI60469	Human polynucleoti

ALIGNMENTS

RESULT 1
AAC03794
ID AAC03794 standard; CDNA; 447 BP.
XX AAC03794:
AC
AC
DT 06-OCT-2000 (first entry)
XX
XX Human secreted protein 5' EST, SEQ ID NO: 3792.
DE
XX Human, 5' EST; expressed sequence tag; secreted protein; CDNA isolation;
KW gene therapy; chromosome mapping; ss.
XX
OS Homo sapiens.
XX
XX EPI033401-A2.
XX
PD 06-SEP-2000.

XX 21-FEB-2000; 2000EP-0200610.
XX
XX 26-FEB-1999; 99US-0122487.
XX
XX (GEST) GENSET.
XX
XX
XX Dumas Mline Edwards J, Duclert A, Giordano J;
XX
XX WPI: 2000-500381/45.
XX
XX P-PSDB; AAG03788.
XX
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
XX obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
XX diagnostic, forensic, gene therapy and chromosome mapping procedures -
XX
XX Claim 1; SEQ ID 3792; 71pp + CD-ROM; English.

The present sequence is one of a large number of 5' ESTs derived from mRNAs encoding secreted proteins. An ORF has been identified within the sequence. The 5' ESTs were prepared from total human RNAs or poly(A) RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of mRNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design expression and secretion vectors.

Alignment scores:	
Pred. No.:	3.61e-20
Score:	193.00
Percent Similarity:	100.00%
Best Local Similarity:	100.00%
Query Match:	100.00%
DB:	21
Length:	447
Matches:	37
Conservative:	0
Mismatches:	0
Indels:	0
Gaps:	0

US-09-513-999C-7869_COPY_1-37 (1-37) x AAC03794 (1-447)

0y 1 MetIlglySerPheAlaLeuGlnAspSerPheSerLeuGlnIleuLeuGlyPro 20
51 ATGGTGCATCTTTGGCTTCAGAGATTCTTTTCACTTTTGCAGGACCTTCCTGGSCCG 110
0y 21 GlnrValLysLeuGlyLeuGlyValCysLeuSerClyCysSerThr 37
111 GAGATGTCGAAACCTCTGGGCTCTCTGTGTGGCTGTAGGGCTGCTCTACT 161
0b

RESULT 2

ID AA242680 standard; cDNA; 447 BP.

AC AA242680;

DT 01-FEB-2000 (first entry)

DE Human 5' EST isolated from a cDNA library SEQ ID NO:439.

KW Human; 5' EST; expressed sequence tag; secreted protein; diagnosis;

KW forensic; location; development; protein synthesis; stability;

XX XX

OS Homo sapiens.

PN W09953051-A2.

PD 21-OCT-1999.

XX 09-APR-1999; 99WO-IB00712.
 XX
 XX 09-APR-1998; 98US-0057719.
 PR 28-APR-1998; 98US-0069047.
 PR
 XX (GEST) GENSET.
 PA
 XX
 XX Dumas Milne Edwards J, Duclert A, Giordano J;
 PI
 XX WPI; 2000-038446/03.
 XX
 DR P-PSDB; AAY65066.
 DR
 XX
 XX Novel secreted protein 5' expressed sequence tag sequences used in
 PT diagnostic, forensic, gene therapy, and chromosome mapping procedures
 XX
 PS Claim 1; Page 402; 837pp; English.

CC AA242265 to AA243075 represent novel 5' expressed sequence tag (EST)
CC sequences corresponding to human secreted proteins. AA56451 to
CC AA565438 represent the EST-related proteins corresponding to AA242265 to
CC AA243052. The 5' ESTs can be used for producing secreted human gene
CC products. They can be used to identify and isolate 5' untranslated
CC regions (UTRs) and upstream regulatory regions which control the
CC location, development stage, rate, and quantity of protein synthesis, as
CC well as stability of mRNA. The ESTs are also useful as probes for
CC chromosome mapping, and to obtain full length cDNA clones. The ESTs can
CC also be used in forensic procedures to identify individuals, or in
CC diagnostic procedures to identify individuals having genetic diseases
CC resulting from abnormal gene expression. The products may also be used in
CC gene therapy protocols. The nucleic acids encoding signal peptides can be
CC used for directing extracellular secretion of a polypeptide or the
CC insertion of a polypeptide into a membrane or importing a polypeptide
CC into a cell. The proteins encoded by the EST sequences may be useful in
CC treating a variety of human conditions. Secreted proteins have
CC therapeutic value, and the identification of new secreted proteins is
CC valuable. AA242249 to AA242264 and AA564544 to AA564550 represent
CC sequences used in the exemplification of the present invention.

Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;

Alignment Scores:	
pred. No.:	3 61e-20
Score:	193.00
Percent Similarity:	100.00%
Best Local Similarity:	100.00%
Query Match:	100.00%
DB:	21
Length:	44
Matches:	37
Conservative:	0
Mismatches:	0
Indels:	0
Gaps:	0

US-09-513-999C-7869_COPY_1_37 (1-37) x AA242680 (1-447)

Oy 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyPro 20
 Db 51 ATGGGTGGACTCTTTGGCTTCAGAGATCTTTTCATCTTGGACGGAGACTTCGGGGCCG 110
 Oy 21 GlutryValIysLeuLeuGlyLeuGlyValCysLeuSerGlyCysSerThr 37
 Db 111 GAGATGATGAAACCTCTGGGCTCTCTGGTGTGGCCGTGAGTGGCTCTCTACT 161

RESULT 3

ID	ABK83461	standard;	CDNA;	128600	BP.

AC ABK83461;

DT 14-AUG-2002 (first entry)

DE Human cDNA differentially expressed in granulocytic cells #32.

KW Human; ss; granulocytic cell; DNA chip; bacterial infection;

KW fungal infection; sterile inflammatory disease; psoriasis;

KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.
 XX
 XX Homo sapiens.
 OS
 PN WO/2002/28999-A2.
 PD 11-APR-2002.
 XX
 PF 03-OCT-2001; 2001MO-US30821.
 XX
 PR 03-OCT-2000; 2000US-237189P.
 XX
 PA (GENE-) GENE LOGIC INC.
 XX
 PI Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
 DR WPI: 2002-435328/46.
 XX
 PT Detecting granulocyte activation by detecting differential expression
 of genes associated with granulocyte activation, which serves as
 diagnostic markers that is useful for monitoring disease states and
 drug toxicity -
 PS
 XX
 PS Claim 1; SEQ ID NO 32; 114pp; English.
 CC
 CC The invention relates to detecting (M1) granulocyte (GC) activation
 (GCA), by detecting the level of expression of gene(s) (Gs) identified by
 CC DNA chip analysis as given in the specification, and comparing
 CC the expression level to an expression level in an unactivated
 CC GC, where differential expression of Gs is indicative of GCA.
 CC Also included are modulating (M2) GA by contacting GC with an agent
 CC that alters the expression of at least one gene in Gs; (2) screening (M3)
 CC for an agent capable of modulating GCA or an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease using the
 CC gene expression profile; (3) detecting (M4) an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease, by detecting the
 CC level of expression in a sample of the tissue of gene(s) from Gs, where
 CC the level of expression of the gene is indicative of inflammation;
 CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,
 CC an allergic response in a subject, exposure of a subject to a pathogen
 CC or sterile inflammatory disease, by contacting a tissue having
 CC inflammation with an agent that modulates the expression of gene(s)
 CC from Gs in the tissue. M1 is useful for detecting GCA, M2 is useful for
 CC modulating GA; M3 is useful for screening an agent capable of modulating
 CC GCA preferably in an inflammation in a tissue; M4 is useful for
 CC detecting an inflammation (especially chronic) in a tissue, an allergic
 CC response in a subject, exposure of a subject to a pathogen or sterile
 CC inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
 CC glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
 CC reperfusion injury, ARDS, adult respiratory distress syndrome,
 CC inflammatory bowel disease, Crohn's disease, ulcerative colitis,
 CC periodontal disease; also bacterial infection, viral infection,
 CC parasitic infection, protozoal infection, fungal infection and M5 is
 CC useful for treating one of the above conditions. The present
 CC sequence represents a gene differentially expressed in granulocytes.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 SO Sequence 128600 BP; 36139 A; 24970 C; 26316 G; 41175 T; 0 other;

Alignment Scores:
 Pred. NO.: 0.0459 Length: 128600
 Score: 95.00 Matches: 21
 Percent Similarity: 64.86% Conservative: 3
 Best Local Similarity: 56.76% Mismatches: 9
 Query Match: 49.22% Indels: 4

```
DB:                               24                                Gaps:                               1
US-09-513-999C-7869_COPY_1_37 (1-37) x ABA63451 (1-128600)
OY      3 GlySerPheAlaLeuGlnAspSerPheSerLeuGln-----gLyLeuLeu 18
        |||:::                ::: |||||
Db 126328 GGGCGTATGTGTGCACCTCCACCCTTGACTTCATTGCAGACGCATTTCGTGGGCTCTG 126269
OY      19 GlYProGluTYrValLysLeuLeuGlyLeuCysValcLysLeuSerGLyCys 35
        ||| ||||||| ||||||| ||||| |||
Db 126268 GGGCGTAGGTATGTAAAGCTCTCGGCTCTGTGCATGCCCTGAGCAGCTGC 126218

RESULT 4
ABA63453/c
ID      ABA63453 standard; DNA; 570 BP.
XX
AC      ABA63453;
XX
DT      01-FEB-2002 (first entry)
XX
DE      Human foetal liver single exon nucleic acid probe #11758.
XX
KW      Human; foetal liver; gene expression; single exon nucleic acid probe; ss
XX
OS      Homo sapiens.
XX
PN      WO200157277-A2.
XX
PD      09-AUG-2001.
XX
PF      30-JAN-2001; 2001WO-US00669.
XX
PR      04-FEB-2000; 2000US-0180312.
PR      26-MAY-2000; 2000US-0207456.
PR      30-JUN-2000; 2000US-0608408.
PR      03-AUG-2000; 2000US-0632366.
PR      21-SEP-2000; 2000US-0234687.
PR      27-SEP-2000; 2000US-0236359.
PR      04-OCT-2000; 2000GB-0024263.
XX
PA      (MOLE-) MOLECULAR DYNAMICS INC.
PI      Penn SG, Hanzel DK, Chen W, Rank DR;
DR      WPI; 2001-483447/52.
XX
PT      Human genome-derived single exon nucleic acid probes useful for
PT      analyzing gene expression in human fetal liver -
PS      Claim 1; SEQ ID NO 11758; 639pp + sequence listing; English.
XX
CC      The invention relates to a single exon nucleic acid probe for
CC      measuring human gene expression in a sample derived from human foetal
CC      liver. The single exon nucleic acid probes may be used for predicting,
CC      measuring and displaying gene expression in samples derived from human
CC      fetal liver. The present sequence is a single exon nucleic acid
CC      probe of the invention.
CC      Note: The sequence data for this patent did not form part of the
CC      printed specification, but was obtained in electronic format directly
CC      from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX
SQ      Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:
Pred. No.:          0.000408                      Length:          570
Score:              88.50                          Matches:           21
Percent Similarity: 61.11%                         Conservative:     1
Best Local Similarity: 58.33%                       Mismatches:       7
Query Match:         45.85%                         Indels:           7
DB:                  22                              Gaps:             1

US-09-513-999C-7869_COPY_1_37 (1-37) x ABA63453 (1-570)
```

```

Oy      1 MetGtGcYSerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyPro 20
Db      568 ATGAATGATATC-----CTGCCTTGCTGGATTCCTTGCGCTG 530
Oy      21 GltTyrValLysLeuLeuGlyLeuCysValCysLeuSerGlyCysSer 36
Db      529 GAGTATGCTAAATTCCTGGCTCTTGCTGCTGCTGCTGAGTGCCGCGCT 482

RESULT 5
ABA30652/c
ID      ABA30652 standard; DNA: 570 BP.
XX
XX      ABA30652;
AC
XX      23-JAN-2002 (first entry)
DT
XX      Probe #9118 for gene expression analysis in human heart cell sample.
DE
XX      Human; gene expression; heart; microarray; vascular system; probe;
XX      cardiovascular disease; hypertension; cardiac arrhythmia;
XX      congenital heart disease; ss.
XX      Homo sapiens.
OS
XX      WO200157274-A2.
XX
XX      09-AUG-2001.
XX
XX      30-JAN-2001: 2001WO-US00666.
XX
XX      04-FEB-2000: 2000US-0180312.
XX      26-MAY-2000: 2000US-0207456.
XX      30-JUN-2000: 2000US-0608408.
XX      03-AUG-2000: 2000US-0632366.
XX      21-SEP-2000: 2000US-0234687.
XX      27-SEP-2000: 2000US-0236359.
XX      04-OCT-2000: 2000GB-0024263.
XX
XX      (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX      Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX      WPI; 2001-48889/53.
XX
XX      Single exon nucleic acid probes for analyzing gene expression in human
XX      hearts -
XX
XX      Claim 1; SEQ ID No 9118; 530bp; English.

The present invention relates to single exon nucleic acid probes for
CC      measuring human gene expression in a sample derived from human heart. The
CC      present sequence is one such probe. The probes may be used for
CC      predicting, measuring and displaying gene expression in samples derived
CC      from the human heart via microarrays. By measuring gene expression, the
CC      probes are useful for predicting, diagnosing, grading, staging,
CC      monitoring and prognosing diseases of the human heart and vascular system
CC      e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
CC      congenital heart disease.
CC      Note: The sequence data for this patent did not form part of the printed
CC      specification, but was obtained in electronic format directly from WIPO
CC      at ftp.wipo.int/pub/published.pct-sequences.
XX
XX      Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:
Pred. NO.:      0.00408      Length:      570
Score:          88.50      Matches:      21
Percent Similarity: 61.11%      Conservative: 1
Best local Similarity: 58.33%      Mismatches: 7
Query Match:     45.85%      Indels:      7
DB:             22      Gaps:        1

US-09-513-999C-7869-COPY_1-37 (1-37) x ABA30652 (1-570)

```

Oy	1	MctgTgcYgYserPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuEngLyPro	20
			:::
Db	568	ATGAATGCATCT-----CCTGCTTGCTGGGATTCCTTGCGCTG	530
Oy	21	GtUrYValLysLeuLeuGlyLeuCysValCysLeuSerGlyCysSer	36
Db	529	GAGTATGTAAATTCCCTGGCTTTGTTGTGTGACCTGAGTCGCCGCTCT	482
RESULT 6			
ID	AAK11985/c		
XX	AAK11985 standard; DNA; 570 BP.		
AC	AAK11985;		
DT	05-NOV-2001 (first entry)		
DX	Human brain expressed single exon probe SEQ ID NO: 11976.		
XX			
KW	Human: brain expressed exon; gene expression analysis; probe:		
KM	microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;		
XX	epilepsy; cancer; SS.		
OS	Homo sapiens.		
PN	WO200157275-A2.		
PD	09-AUG-2001.		
PF	30-JAN-2001; 2001WO-US00667.		
PR	04-FEB-2000; 2000US-0180312.		
PR	26-MAY-2000; 2000US-0207456.		
PR	30-JUN-2000; 2000US-0608408.		
PR	03-AUG-2000; 2000US-0632366.		
PR	21-SEP-2000; 2000US-0234687.		
PR	27-SEP-2000; 2000US-0236359.		
PR	04-OCT-2000; 2000GB-0024263.		
PA	(MOLE-) MOLECULAR DYNAMICS INC.		
PI	Penn SG, Hanzel DK, Chen W, Rank DR;		
DR	WPI; 2001-483446/52.		
PT	Single exon nucleic acid probes for analyzing gene expression in human brains -		
PS	Example 4; SEQ ID NO: 11976; 650bp + Sequence listing; English.		
CC	The present invention provides a number of single exon nucleic acid		
CC	probes which are derived from genomic sequences expressed in the human		
CC	brain. They can be used to measure gene expression in brain cell samples		
CC	which may enable the diagnosis and improved treatment of nervous system		
CC	diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,		
CC	epilepsy and cancers. The present sequence is one of the probes of the		
CC	invention.		
SQ	Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;		
Alignment Scores:			
Pred. NO.:	0.000408	Length:	570
Score:	88.50	Matches:	21
Percent Similarity:	61.11%	Conservative:	1
Best Local Similarity:	58.33%	Mismatches:	7
Query Match:	45.85%	Indels:	7
DB:	22	Gaps:	1
Oy	1	MctgTgcYgYserPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuEngLyPro	20
			:::
Db	568	ATGAATGCATCT-----CCTGCTTGCTGGGATTCCTTGCGCTG	530

OY 21 GluTyrValIysLeuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
|||||
Db 529 GAGTATGTAATAATTCCTGGGCTTTGTGTGTGCTGCGCTGAGTGGCGGCTCT 482

RESULT 7

AAK37688/c
ID AAK37688 standard; DNA; 570 BP.

AC AAK37688;

DT 06-NOV-2001 (first entry)

DE Human bone marrow expressed single exon probe SEQ ID NO: 12245.

KW Human; bone marrow expressed exon; gene expression analysis; probe;
microarray; cancer; leukemia; lymphoma; myeloma; ss.

OS Homo sapiens.

WO200157276-A2.

PD 09-AUG-2001.

PE 30-JAN-2001; 2001WO-US00668.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

PA (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

PT WPI: 2001-488900/53.

PS Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human bone marrow -

XX Example 4; SEQ ID NO: 12245; 658bp + Sequence Listing; English.

CC The present invention provides a number of single exon nucleic acid

CC probes which are derived from genomic sequences expressed in the human

CC bone marrow. They can be used to measure gene expression in bone marrow

CC samples, which may enable the improved diagnosis and treatment of cancers

CC such as lymphoma, leukemia and myeloma. The present sequence is one of

CC the probes of the invention.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:

Pred. No.: 0.000408 Length: 570

Score: 88.50 Matches: 21

Percent Similarity: 61.11% Conservative: 1

Best Local Similarity: 58.33% Mismatches: 7

Query Match: 45.85% Indels: 7

DB: Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x AAK37688 (1-570)

OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyPro 20

Db 568 ATGAATGATCT-----CCTGCTTCTGGGATTCTTGGGCTG 530

OY 21 GluTyrValIysLeuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36

Db 529 GAGTATGTAATAATTCCTGGGCTTTGTGTGTGCTGCGGCTCT 482

RESULT 8

AA118447/c
ID AA118447 standard; DNA; 570 BP.

AC AA118447;

DT 12-OCT-2001 (first entry)

DE Probe #8380 for gene expression analysis in human cervical cell sample.

KW Probe; human; microarray; gene expression; cervical epithelial cell;

KW cervical cancer; ss.

OS Homo sapiens.

WO200157278-A2.

PD 09-AUG-2001.

PE 30-JAN-2001; 2001WO-US00670.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

PA (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

PT WPI: 2001-488901/53.

PS Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human cervical epithelial cells -

XX Claim 25; SEQ ID No 8380; 487bp; English.

CC The present invention relates to human single exon nucleic acid probes

CC (SNP). The present sequence is one such probe. The SNPs are derived

CC from human HeLa cells. The SNPs can be used to produce a single exon

CC microarray, which can be used for measuring human gene expression in a

CC sample derived from human cervical epithelial cells. By measuring gene

CC expression, the probes are therefore useful in grading and/or staging

CC of diseases of the cervix, notably cervical cancer.

CC Note: The sequence data for this patent did not form part of the printed

CC specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:

Pred. No.: 0.000408 Length: 570

Score: 88.50 Matches: 21

Percent Similarity: 61.11% Conservative: 1

Best Local Similarity: 58.33% Mismatches: 7

Query Match: 45.85% Indels: 7

DB: Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x AA118447 (1-570)

OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyPro 20

Db 568 ATGAATGATCT-----CCTGCTTCTGGGATTCTTGGGCTG 530

OY 21 GluTyrValIysLeuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36

Db 529 GAGTATGTAATAATTCCTGGGCTTTGTGTGTGCTGCGGCTCT 482

RESULT 9

AA143563/c
ID AA143563 standard; DNA; 570 BP.

```

XX AA143563;
AC 17-OCT-2001 (first entry)
DT
DE Probe #12249 used to measure gene expression in human placenta sample.
DE
KW Probe; microarray; human; placenta; antenatal diagnosis;
KW genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200157272-A2.
XX
PD 09-AUG-2001.
XX
PE 30-JAN-2001; 2001WO-US00663.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024253.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI: 2001-488897/53.
XX
PS Human genome-derived single exon nucleic acid probes useful for
PS analyzing gene expression in human placenta -
XX
PT Claim 25; SEQ ID No 12249; 654pp; English.
XX
PS
XX The present invention relates to single exon nucleic acid probes (SENP).
XX The present sequence is one such probe. The probes are useful for
XX producing a microarray for predicting, measuring and displaying gene
XX expression in samples derived from human placenta. The probes are useful
XX for antenatal diagnosis of human genetic disorders.
XX
XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
XX
XX Alignment Scores:
XX Pred. No.: 0.000408 Length: 570
XX Pre: 88.50 Matches: 21
XX Best Similarity: 61.11% Conservative: 1
XX Best Local Similarity: 58.33% Mismatches: 7
XX Query Match: 45.85% Indels: 7
XX DB: Gaps: 1
XX
XX
XX US-09-513-999C-7869_COPY_1_37 (1-37) x AA143563 (1-570)
XX
XX
XX 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyPro 20
XX ||| |||||
XX Db 568 ATGATGATCT-----CCTGCTTGGCGGATTCCTGGGCTG 530
XX
XX
XX 21 GluIyrValIysLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
XX ||||| ||||| ||||| ||||| |||||
XX Db 529 GAGTATGTAATAATTCCTGGGTCTTGTGTGCTGATCGGCGCTCT 482
XX
XX
XX RESULT 10
XX ID ABS11680 standard; DNA: 570 BP.
XX
XX ABS11680;
XX AC
XX 19-AUG-2002 (first entry)
XX
XX Human genome-derived single exon probe from lung SEQ ID No 11671.
XX

```

Human; stinagle exon probe; asthma; lung cancer; COPD; ILD; chronic obstructive pulmonary disease; interstitial lung disease; familial idiopathic pulmonary fibrosis; neurofibromatosis; tuberous sclerosis; Gaucher's disease; Niemann-Pick disease; Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis; pulmonary histiocytosis; lymphangioleiomyomatosis; Kargener syndrome; pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia; primary ciliary dyskinesia; pulmonary hypertension; hyaline membrane disease.

Homo sapiens.

WO200186003-A2.

15-NOV-2001.

30-JAN-2001; 2001WO-US00665.

04-FEB-2000; 2000US-180313P.

26-MAY-2000; 2000US-20745P.

30-JUN-2000; 2000US-0608408.

03-AUG-2000; 2000US-063236P.

21-SEP-2000; 2000US-234687P.

27-SEP-2000; 2000US-236359P.

04-OCT-2000; 2000GB-0024253.

(MOLE-) MOLECULAR DYNAMICS INC.

Penn SG, Hanzel DK, Chen W, Rank DR; WPI; 2002-114183/15.

Spatially-addressable set of single exon nucleic acid probes, used to measure gene expression in human lung samples -

Claim 1; SEQ ID No 11671; 634pp; English.

The invention relates to a spatially-addressable set of single exon nucleic acid probes for measuring gene expression in a sample derived from human lung comprising single exon nucleic acid probes having one of 12614 nucleic acid sequences mentioned in the specification, or their complements or the 12387 open reading frames derived from the 12614 probes. Also included are a microarray comprising the novel set of probes, the novel set of probes which hybridise at high stringency to a nucleic acid expressed in the human lung; measuring gene expression in a sample derived from human lung, comprising (a) contacting the array with mRNA, and (b) measuring the label detectably bound to each probe of the array; identifying exons in a eukaryotic genome, comprising (a) algorithmically predicting at least one exon from genomic sequences of the eukaryote; and (b) detecting specific hybridisation of detectably labeled nucleic acids from eukaryote lung mRNA, to a single exon probe, having a fragment identical to the predicted exon, the probe is included in the above mentioned microarray; assigning exons to a single gene, comprising (a) identifying exons from genomic sequence by the method above and (b) measuring the expression of each of the exons in several tissues and/or cell types using hybridisation to a single exon microarrays having a probe with the exon, where a common pattern of expression of the exons in the tissues and/or cell types indicates that the exons should be assigned to a single gene; a peptide comprising one of 12011 sequences, mentioned in the specification, or encoded by the genes of 12011 sequences, mentioned in the specification, and for gene probes/open reading frames (ORF). The probes are used for gene expression analysis, and for identifying exons in a gene, particularly using human lung derived mRNA and for the study of lung diseases such as asthma, lung cancer, chronic obstructive pulmonary disease (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease, Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis, pulmonary alveolar proteinosis, Kargener syndrome, fibrocystic pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension and hyaline membrane disease. The present sequence is a single exon probe of the invention.

CC Note: The sequence data for this patent did not form part
CC of the printed specification, but was obtained in electronic
CC format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX

SO Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:

Pred. No.:	Length:	Score:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
0.000408	570	88.50	21	1	7	7	1

Percent Similarity: 61.11%
Best Local Similarity: 58.33%
Query Match: 45.85%

DB: 24

US-09-513-999c-7869_COPY_1_37 (1-37) x ABS11680 (1-570)

OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyPro 20
|||
568 ATGAATGATCTC-----CCTGCCTTCCTGGATTCCTTGGCTG 530
|||
21 GUTYrVAllyLsLeuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
|||||
529 GAGTATGTAATAATCTCTGCGCTCTTGTGTGTCTGCTGAGTGGCCGCTCT 482

RESULT 11
AAC68089/c
ID AAC68089 standard; cDNA; 1982 BP.
XX
XX AAC68089;
AC
XX
XX 20-FEB-2001 (first entry)
DE Human secreted protein cDNA sequence #9.
XX
XX Cytostatic; immunosuppressive; neurotropic; neuroprotective; antiviral;
XX antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;
XX vulnerability; anticonvulsant; antibacterial; antifungal; antiparasitic;
XX cardiac; gene therapy; cancer; immune disorder; cardiovascular disorder;
XX neurological disease; infection; human; secreted protein; ss.
OS
XX Homo sapiens.
XX
XX WO200058335-A1.
PN
XX 05-OCT-2000.
PD
XX 22-MAR-2000; 2000WO-US07534.
PF

26-MAR-1999; 99US-0126598.
22-DEC-1999; 99US-0171504.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX (ROSE/) ROSEN C A.
PA
XX Rosen CA, Ruben SM, Komatsuolis G;
XX
XX WPI: 2000-611702/58.
XX P-PSDB; AAB37356.
DR
XX
XX Nucleic acids encoding human secreted proteins, used to treat, prevent,
XX ameliorate or diagnose conditions such as cancer, and autoimmune
XX diseases e.g. arthritis -
PT
XX
XX Claim 1; Pages 321-322; 387pp; English.

The invention relates to the isolation of genes AAC68081-C68127 encoding
47 human secreted proteins AAB37348-B37394. The genes can be used to
generate fusion proteins by linking to the gene for the human
immunoglobulin G Fc portion (AAC68072) for increasing the stability of
the fusion protein as compared to the human protein only. The genes and
proteins are useful for preventing, ameliorating or treating medical
conditions, e.g. by protein or gene therapy. The genes are isolated

CC from a range of human tissues disclosed in the specification. The
CC nucleic acids, proteins, antibodies and (anti)agonists are useful in
CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
CC and ovarian cancer, and other cancers of the adrenal gland, bone, bone
CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
CC colitis; (c) cardiovascular disorders such as myocardial ischaemia; (d)
CC wound healing; (e) neurological diseases e.g. cerebral anoxia and
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
CC and parasitic infections.
XX

SO Sequence 1982 BP; 549 A; 513 C; 471 G; 448 T; 1 other;

Alignment Scores:

Pred. No.:	Length:	Score:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
0.00243	1982	88.00	20	1	10	0	0

Percent Similarity: 67.74%
Best Local Similarity: 64.52%
Query Match: 45.60%

DB: 21

US-09-513-999c-7869_COPY_1_37 (1-37) x AAC68089 (1-1982)

OY 6 AlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGlyrVAllyLsLeu 25
|||||
Db 1156 GCTTTCAGAGATTGACAGTACTTTCTGCGAACCAGCGCGAGTATGTAAGCTG 1097
|||||
OY 26 LeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
|||||
Db 1096 CTGGGCTCTGACAGTGTCTCAAGCGGCTACTCT 1064

RESULT 12
AAS91827/c
ID AAS91827 standard; cDNA; 1299 BP.
XX
XX AAS91827;
AC
XX
XX 13-FEB-2002 (first entry)
DE
XX
XX DNA encoding novel human diagnostic protein #27631.
XX
XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX Homo sapiens.
XX
XX WO200175067-A2.
PN
XX 11-OCT-2001.
PD
XX 30-MAR-2001; 2001WO-US08631.
PF

31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Yang YF;
XX
XX WPI: 2001-639362/73.
XX P-PSDB; ABG27640.
DR
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -
PT
XX
XX Claim 1; SEQ ID No 27631; 103pp; English.

The invention relates to isolated polynucleotide (I) and
polypeptide (II) sequences. (I) is useful as hybridisation probes,

PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
XX
P-PSDB: ABG24247.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 1: SEQ ID No 24238; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS6197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ Sequence 1909 BP; 705 A; 443 C; 389 G; 372 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.0134 Length: 1909
Score: 83.00 Matches: 18
Percent Similarity: 60.61% Conservative: 2
Best Local Similarity: 54.55% Mismatches: 13
Query Match: 43.01% Indels: 0
DB: Gaps: 0
XX
09-513-999C-7869_COPY_1_37 (1-37) x AAS88434 (1-1909)
XX
QY 4 SerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrVal 23
DB 205 TCAAGACCTTCCTGAGTACGATCCATTCCTGCGAGGACCGAGGACGAGTACATA 146
XX
QY 24 LysLeuGlyLeuGlyValCysLeuSerGlyCysSer 36
DB 145 AAGCCCTGGTCTTGTGATGCTGCTGAGACAGCTCTCC 107
XX
RESULT 15
ID AAH98980
AAH98980 standard: cDNA; 609 BP.
XX
AC AAH98980;
XX
DT 12-OCT-2001 (first entry)
XX
DE Human EST-derived coding sequence SEQ ID NO: 837.
XX
XX Human: sheep; pig; cow; fruit fly; yeast; hamster; macaque; horse;
KW tomato; monkey; dog; sea urchin; expressed sequence tag; EST;
KW diagnostics; forensic test; gene mapping; genetic disorder;
KW biodiversity; gene therapy; nutrition; ss.

XX
OS Homo sapiens.
XX
PN WO200154477-A2.
XX
XX
PD 02-AUG-2001.
XX
XX 25-JAN-2001; 2001WO-US02687.
XX
PF 25-JAN-2000; 2000US-0491404.
PR 17-JUL-2000; 2000US-0617746.
PR 03-AUG-2000; 2000US-0631451.
PR 15-SEP-2000; 2000US-0663870.
XX
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;
PI Cao Y, Drmanac RA, Zhang J, Werhman T;
XX
XX WPI: 2001-476164/51.
DR P-PSDB: AAM24321.
XX
PT Isolated polypeptide for treatment of diseases, diagnostics, raising
PT antibodies and research use -
XX
PS Claim 1: Page 694; 1275bp; English.
XX
XX The present invention provides the protein and coding sequences of novel
CC proteins from a variety of organisms, including human, dog, cat, horse,
CC cow, pig, hamster, monkey, macaque, yeast, bacteria, fruit fly, sea
CC urchin and tomato. These were derived from expressed sequence tags (ESTs)
CC from the organism of interest. They can be used in diagnostics,
CC forensics, gene mapping, identification of mutations, to assess
CC biodiversity and for nutritional purposes. The present sequence is a cDNA
CC of the invention.
XX
SQ Sequence 609 BP; 99 A; 162 C; 173 G; 175 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.0176 Length: 609
Score: 78.00 Matches: 17
Percent Similarity: 71.43% Conservative: 3
Best Local Similarity: 60.71% Mismatches: 8
Query Match: 40.41% Indels: 0
DB: Gaps: 0
XX
US-09-513-999C-7869_COPY_1_37 (1-37) x AAH98980 (1-609)
XX
QY 6 AlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeu 25
DB 273 GCCCTGCCGAGATCCATCCATTCCTTGGGATCCTAAGCTGAGTATGTAAGCTC 332
XX
QY 26 LeuGlyLeuGlyValCysLeuSer 33
DB 333 CTGGCTCTCTATGTCCTGAGC 356
XX
Search completed: April 24, 2003, 23:01:50
Job time : 165 secs

GenCore version 5.1.4_p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: April 24, 2003, 22:34:13 ; Search time 1054 Seconds

(without alignments)
568.532 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37

Perfect score: 193

Sequence: 1 MGSEFALQDSRSSLOGILGPEYVKILGICVCLSGCSTR 37

Scoring table:

BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Trashed: 16154066 segs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:
-MODEL=frame+ p2n.model -DEV=xlh
-Q=/cgn2.1/USPQC_spool/US09513999/unat_18042003.170937.28370/app_query.fasta.1.199
-DB=EST -QFMT=fastap -SUFFIX=p2n.rst -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blsum62 -TRANS=human40.cdi -LIST=45
-DOCALLIGN=200 -THR.SCORE=pct -THR.MAX=100 -THR.MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pct -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999 @cgn.1.1456 @unat.18042003.170937.28370 -NCPU=6 -ICPU=3
-NO_XLPHY -NO_MMAP -LARGESTUEVY -NEG.SCORES=0 -MATT -LONGLOG -DEV.TIMEOUT=120
-WARN.TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

EST :
1: em_estba:*
2: em_esthum:*
3: em_estlin:*
4: em_estlun:*
5: em_estlov:*
6: em_estlpl:*
7: em_estlro:*
8: em_hlcc:*
9: gb_estcl:*
10: gb_est2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rtd:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	193	100.0	470	17	A0770688	A0770688 HS.5368_B
2	169	87.6	525	17	A0165256	A0165256 HS.3025_B
3	129	66.8	864	17	A0739614	A0739614 HS.3505_A
4	109	56.5	519	17	A0186743	A0186743 HS.3109_B
5	109	56.5	533	17	A0358845	A0358845 HS.5028_B
6	108	56.0	628	17	A0237815	A0237815 RPT-11-70
7	105	54.4	292	17	A0508480	A0508480 RPT-11-2
8	105	54.4	632	17	AG146082	AG146082 Pan trogl
9	104	53.9	666	17	A0427698	A0427698 C17B1-E1
10	102	52.8	418	9	AA493535	AA493535 ng75g10.s
11	102	52.8	556	17	A0384817	A0384817 RPT-11-13
12	102	52.8	695	17	AG179297	AG179297 Pan trogl
13	101	52.3	703	17	A0534396	A0534396 RPT-11-3
14	100	51.8	634	17	AG160901	AG160901 Pan trogl
15	100	51.8	639	17	AG151043	AG151043 Pan trogl
16	100	51.8	676	17	AG061401	AG061401 Pan trogl
17	100	51.8	769	17	A0899390	A0899390 HS.5234_A
18	99	51.3	362	17	A0102366	A0102366 HS.3040_A
19	99	51.3	410	17	A0442274	A0442274 HS.5137_A
20	99	51.3	416	17	A0182486	A0182486 HS.3077_A
21	99	51.3	453	17	A0437684	A0437684 HS.5137_A
22	99	51.3	485	17	A0671849	A0671849 HS.5462_A
23	99	51.3	635	17	A0390599	A0390599 C17B1-E1
24	98	50.8	546	17	A0435071	A0435071 HS.5114_B
25	98	50.8	553	17	A2521751	A2521751 RPT-11-1
26	97	50.3	597	17	A0506884	A0506884 RPT-11-3
27	96.5	50.0	410	17	A0531865	A0531865 RPT-11-3
28	96.5	50.0	672	17	A0551272	A0551272 RPT-11-4
29	96	49.7	347	17	A0631315	A0631315 RPT-11-4
30	96	49.7	444	17	A0463109	A0463109 HS.5211_A
31	96	49.7	663	17	AG091225	AG091225 Pan trogl
32	95	49.2	380	17	A0207172	A0207172 HS.3239_B
33	95	49.2	519	17	A0139984	A0139984 HS.5106_A
34	95	49.2	546	17	A0541696	A0541696 RPT-11-3
35	95	49.2	672	17	AG051939	AG051939 Pan trogl
36	94	48.7	363	17	A0120796	A0120796 HS.3076_A
37	94	48.7	425	17	A0683450	A0683450 HS.5432_B
38	94	48.7	452	17	A0534129	A0534129 RPT-11-3
39	94	48.7	516	17	A0457001	A0457001 HS.5151_A
40	94	48.7	546	17	A0333597	A0333597 HS.5008_A
41	93	48.2	529	17	A0881246	A0881246 HS.5137_B
42	92	47.7	580	17	A0532835	A0532835 RPT-11-3
43	91	47.2	463	17	A0550196	A0550196 RPT-11-4
44	91	47.2	546	17	A0468922	A0468922 HS.5139_A
45	91	47.2	559	17	A0385332	A0385332 RPT-11-14

ALIGNMENTS

RESULT 1
A0770688
LOCUS
DEFINITION
A0770688
ACCESSION
A0770688
VERSION
A0770688
KEYWORDS
SOURCE
ORGANISM
human.
GSS.
A0770688 470 bp DNA linear GSS 28-JUL-1999
HS.5368.B2.C08.SP6E RPT-11 Human Male BAC Library Homo sapiens
genomic clone Plate=944 COL-16 Row=F, DNA sequence.

REFERENCE
AUTHORS
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Mahatras,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
1 (bases 1 to 470)
Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.
Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and

TITLE	Hood, L.
JOURNAL	Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
MEDLINE	Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
COMMENT	99380589 Contact: Mahairas GG, Wallace JC, Hood L

Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahatrais GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPOT-11. For BAC library availability, please contact Plietser de Jong (plietserdejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm) or from Research Genetics (<http://www.htsc.washington.edu>)
<http://www.htsc.washington.edu>
Plate: 944 row: F column: 16
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 470.

FEATURES
source

Location/Qualifiers
1. .470

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_11b="RPC1-11 Human Male BAC Library"
/sex="male"
/notes="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI methylase. Size selected DNA was cloned into the pBACe3.6 vector at EcoRI sites"
BASE COUNT      83 a      112 c      131 g      141 t      3 others

```

ALIGNMENT	ORIGIN
Alignment Scores:	
Pred. No.:	4,59e-16
Score:	193.00
Percent Similarity:	100.00%
Best Local Similarity:	100.00%
Query Match:	100.00%
DB:	17
US-09-513-999C-7869-COPY_1_37 (1-37) x AQ70688 (1-470)	
Length:	470
Matches:	37
Conservative:	0
Mismatches:	0
Indels:	0
Gaps:	0

1 MetGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyPro 20
 103 ATGGGTGATCTTTTGCCTTCACAGATTCCTTTTCATCTTTGCAGGAGCTCTGGGGCGG 162
 21 GlnTyrValLysLeuLeuGlyLeuGlyValCysLeuSerGlyCysSerThr 37
 163 GAGTATGTAACACCTCTGGGCTCTGTGTGTGGCCGATGGCTGCTACT 213

RESULT 2	525 bp	DNA	linear	GSS 16-OCT-1998
LOCUS	HS-3025_B2.G06_T7	CIT	Approved Human Genomic Specim	Library D Homo
DEFINITION	sapiens genomic clone Plate-3025 Col-12 Row=N,		DNA sequence.	
ACCESSION	AO165256			
VERSION	AO165256.1	GI:3563451		
KEYWORDS	GSS.			
SOURCE	human.			
ORGANISM	Homo sapiens			

REFERENCE
Eukaryota: Metazoa: Chordata: Caninata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Primates: Catarrhini: Homnidae: Homo. 1 (bases 1 to 525)
Mahairas,G.C., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Kellier,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
TITLE
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL
MEDLINE
99380589
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3025 row: N column: 12
Class: BAC ends
High quality sequence s.top: 525.

FEATURES

Location/Qualifiers
1. .525

BASE COUNT	ORIGIN
102 a	137 g 143 t 4 others
	E-Coli DH10b"
	/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in
	/sex="male"
	/clone.lib="CIR Approved Human Genomic Sperm Library D
	/clone="Plate3025 COL-12 Row=N"
	/db_xref="taxon:9606"
	/organism="Homo sapiens"

Alignment Scores:

Pred. No.:	8.15e-13	Length:	525
Score:	169.00	Matches:	33
Percent Similarity:	91.89%	Conservative:	1
Best Local Similarity:	89.19%	Mismatches:	3
Query Match:	87.56%	Indels:	0
DB:	17	Gaps:	0

US-09-513-999C-7869_COPY_1_37 (1-37) x A0165256 (1-525)

QY 1 MetcylglyserPhenalaleuglnaspserPheaserSerleuglnglyLeuleuglyPro 20
| | | | | : | | | | |
Db 66 ATGGGCGGATCTTTCCCTTCAGAGAATAATTATTCATCTTTTGCACGGACTTTGTGGGGCCG 12

Oy 21 GufyValIysIeuleuclIyIeucyValIcysIeuseuSergIyCysserThr 37
 |||||
 Db 126 GAGTATGTAAACCTCCTCCTGCTCTGTGTGTGCTGAGTGGCTGCTGCTACT 176

RESULT 3

LOCUS	864 bp	DNA	linear	GSS 16-JUL
DEFINITION	HS-5505_A1_A09_T7A RPCI-11 Human Male BAC library Homo sapiens genomic clone Plate=1081 Col=17 Row=A, DNA sequence.			

ACCESSION	AQ739814
VERSION	AQ739814.1
	GI:5517336

KEYWORDS
SOURCE
ORGANISM

REFERENCE
Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo. (bases 1 to 854)

AUTHORS

G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and

TITLE Hood, L.
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

Journal of the
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589

COMMENT

Mahairas GG, Wallace JC, Hood L

University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPc1-11. For BAC library availability, please contact Pieter de Jong (Pieter@dejong.med.buffalo.edu) Clones may be purchased from

BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (<http://www.htsc.washington.edu>)
Plate: 1081 row: A column: 17
Seq primer: T7
Class: BAC ends
High quality sequence stop: 864.

FEATURES

Location/Qualifiers
1. 864

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=1081 Col=17 Row=A"
/clone_lib="RPC1-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site 1: EcoRI; Site 2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBAC3.6 vector at EcoRI sites"

BASE COUNT

247 a 182 c 178 g 257 t

IN

Alignment Scores:

Prod. No.: 3.03e-07 Length: 864
Score: 129.00 Matches: 23
Percent Similarity: 92.86% Conservative: 3
Best Local Similarity: 82.14% Mismatches: 2
Query Match: 66.84% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AQ739814 (1-864)

OY 9 AspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeuGlyLeu 28
||| |||||:||||| |||||:|||||:|||||:|||||:|||||:|||||
Db 585 GACAAATTTCCACCTTGGCAGGGGCTCGGGTCCACAGTATGTAAACTCTGGGCTC 526

OY 29 CysValCysLeuSerGlyCysSer 36
|||||:|||||:|||||:|||||:|||||:|||||:|||||

Db 525 TCGGTGTGCTGAGTGGCTGCTCT 502

RESULT 4

LOCUS AQ186743/c

DEFINITION HS_3109_B1_A06_T7 CTF Approved Human Genomic Sperm Library D Homo

Accession sapiens genomic clone Plate=3109 Col=11 Row=B, DNA sequence.

VERSION AQ186743

KEYWORDS GSS.

ORGANISM

human.

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 3109 row: B column: 11

Class: BAC ends

High quality sequence stop: 519.

Location/Qualifiers

1. 519

source

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3109 Col=11 Row=B"
/clone_lib="CTF Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT

144 a 155 c 122 g 93 t 5 others

ORIGIN

Alignment Scores:

Prod. No.: 7.79e-05 Length: 519
Score: 109.00 Matches: 21
Percent Similarity: 82.76% Conservative: 3
Best Local Similarity: 72.41% Mismatches: 5
Query Match: 56.48% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AQ186743 (1-519)

OY 8 GlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeuGly 27
:||||| |||||:||||| |||||:|||||:|||||:|||||:|||||

Db 399 AAGGATGGATCTTCTCCCTGGCGGGAATCCTGGGCTGAGATGTAAACTCTGGGT 340

OY 28 LeuCysValCysLeuSerGlyCysSer 36
|||||:|||||:|||||:|||||:|||||:|||||:|||||

Db 339 CTCTGTGTGCTGCCAGTGGCTGCTCT 313

RESULT 5

LOCUS AQ358845

DEFINITION HS_5028_B1_B02_T7 RPC11 Human Male BAC Library Homo sapiens

Accession genomic clone Plate=604 Col=3 Row=D, DNA sequence.

VERSION AQ358845.1 GI:4207721

KEYWORDS GSS.

SOURCE

human.

ORGANISM

human.

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Clones may be purchased from Research Genetics (<http://www.htsc.washington.edu>)

BAC end Web Server: <http://www.htsc.washington.edu>

Plate: 604 row: D column: 3

Seq primer: T7

Class: BAC ends

High quality sequence stop: 533.

Location/Qualifiers

1. 533

source

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate=604 Col=3 Row=D"

/clone_lib="RPC11 Human Male BAC Library"

/sex="male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; RPC11 Human Male BAC Library"

BASE COUNT

98 a 132 c 139 g 151 t 13 others

ORIGIN

VERSION	A0427698.1	GI:4500605
KEYWORDS	GSS.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.	
AUTHORS	1 (bases 1 to 666) Zhao,S., Adams,M.D., Nierman,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.	
TITLE	Use of BAC End Sequences from Caltech Libraries for Sequence-Ready Map Building Unpublished (1997)	
JOURNAL	Other_GSSS: CITB1-EI-2575E9.TF	
COMMENT	Contact: Shaying Zhao, William Nierman, Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel.: 301 838 0200 Fax: 301 838 0208 Email: hbe@tigr.org Clones are available from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends	
FEATURES	Location/Qualifiers	
Source	1..666 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_lib="2575E9" /clone_lib="CITB1-EI" /sex="male" /cell_type="sperm" /note="Vector: pBelBAC11; Site_1: EcoRI; Site_2: EcoRI; Caltech Human BAC Library D"	
BASE COUNT	136 a 157 c 171 g 201 t 1 others	
ORIGIN		
Alignment Scores:		
Pred. NO.:	0.000479	Length: 666
Score:	104.00	Matches: 22
Percent Similarity:	76.47%	Conservative: 4
Best local Similarity:	64.71%	Mismatches: 8
Query Match:	53.89%	Indels: 0
DB:	17 gaps:	0
US-09-513-999C-7869-COPY_1.37 (1-37) x A0427698 (1-666)		
OY	3 GlycerPhaealalauglInaspSerPheSerSerLeuInglyLeuLengLYProglutyr 22 ::: : ::::: ::: Db 365 GGACTTCAGCGCCCTGGCTGTGATTCGACGAAGTTTGTGGGATCCTTGCGGCAGACTTT 424	
OY	23 VallysLeuEnglyLeuCysValCysLeuSerclgycySer 36 Db 425 GTAAGCTCCTGGCTGTGTGTGTGACTGACGAGCTGCTCT 466	
RESULT 10		
LOCUS	AA493535/c	
DEFINITION	AA493535 418 bp mRNA EST 18-AUG-1997	
ACCSSION	ng75910.s1 NC1_CGAP_Pf6 Homo sapiens CDNA clone IMAGE:940674	
VERSION	AA493535	
KEYWORDS	sequence. similar to contains element PTR/ repetitive element ; , mRNA	
SOURCE	EST . human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homniidae; Homo.	
AUTHORS	1 (bases 1 to 418) NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap .	
TITLE	National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index	

PRIMERS
Sequencing: T7
LIBRARY
Vector : pBAC3.6
R.Site 1 : ECORI
R.Site 2 : EcoRI.

FEATURES
Source
1. .695
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="RP43-051111.TJ"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RPCI-43 Chimpanzee Male BAC Library"

BASE COUNT 243 a 135 c 122 g 192 t
ORIGIN
3 others

Alignment Scores:

Pred. No.: 0.000929 Length: 695
Score: 102.00 Matches: 19
Percent Similarity: 78.57% Conservative: 3
Local Similarity: 67.86% Mismatches: 6
Query Match: 52.85% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AG179297 (1-695)

OY 9 AsperPheserSerLeuGlnGlyLeuLeuGlyProGluTyrValLysLeuGlyLeu 28

DB 135 GACGATATCCCTCCAGGATCATGCGCCAGATATGTAATAAATTCCTGGGTCTC 194

OY 29 CysValCysLeuSerGlyCysSer 36

DB 195 TGTGTATCTGATGAGTGGCTGCTCT 218

RESULT 13
AO534396 703 bp DNA linear GSS 18-MAY-1999
LOCUS
RPCI-11-38013.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-38013,
DEFINITION
DNA sequence.

ACCESSION
AO534396
VERSION
AO534396.1 GI:4846086

KEYWORDS
GSS.
SOURCE
human.

ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE
1 (bases 1 to 703)
Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter
,J.C.
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
Unpublished (1997)

JOURNAL
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org

COMMENT
Clones are derived from the human BAC Library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@jng.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
Research Genet cs (info@resgen.com). BAC end search page:
http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
Seq primer: SP6
Class: BAC ends.

FEATURES
Source
1. .703
/organism="Homo sapiens"
/db_xref="GDB:7645730"
/db_xref="taxon:9606"

/clone="RPCI-11-38013"
/clone_lib="RPCI-11"
/sex="Male"
/cell_type="lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC Library"

BASE COUNT 201 a 194 c 165 g 143 t
ORIGIN
143 t

Alignment Scores:

Pred. No.: 0.00128 Length: 703
Score: 101.00 Matches: 17
Percent Similarity: 75.00% Conservative: 4
Best Local Similarity: 60.71% Mismatches: 7
Query Match: 52.33% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AO534396 (1-703)

OY 9 AsperPheserSerLeuGlnGlyLeuLeuGlyProGluTyrValLysLeuGlyLeu 28

DB 389 GATGATATCCCTCCAGGATCATGCGCCAGATATGTAATAAATTCCTGGGTCTC 330

OY 29 CysValCysLeuSerGlyCysSer 36

DB 329 TGTATGTCTGATGAGTGGCTGCTCT 306

RESULT 14
AG160901 634 bp DNA linear GSS 09-JAN-2002
LOCUS
Pan troglodytes DNA, clone: RP43-026N23.T7, genomic survey
DEFINITION
sequence.

ACCESSION
AG160901
VERSION
AG160901.1 GI:16690579

KEYWORDS
GSS.
SOURCE
Male BAC Library clone:RP43-026N23.T7.
Pan troglodytes

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.

REFERENCE
1
Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
TotoKI,Y., Watanabe,H. and Sakaki,Y.
BAC end sequences of Library RPCI-43
Unpublished
2 (bases 1 to 634)
TotoKI,Y., Watanabe,H. and Sakaki,Y.
Direct Submission
Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-chou,Tsuri-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail:chimbse@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the chimpanzee BAC library RPCI-43 This BAC
end was generated during the R&D process and may have higher chance
of clone tracking errors.

COMMENT
Clones are derived from the chimpanzee BAC library RPCI-43 This BAC
end was generated during the R&D process and may have higher chance
of clone tracking errors.

FEATURES
Source
1. .634
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="RP43-026N23.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RPCI-43 Chimpanzee Male BAC Library"

BASE COUNT 116 a 168 c 156 g 194 t
ORIGIN

Alignment Scores:

Pred. No.: 0.00154 Length: 634
 Score: 100.00 Matches: 19
 Percent Similarity: 82.14% Conservative: 4
 Best Local Similarity: 67.86% Mismatches: 5
 Query Match: 51.81% Indels: 0
 DB: 17 Gaps: 0

US-09-513-999c_COPY_1_37 (1-37) x AG151043 (1-659)

QY 9 AspserPheSerLeuGlnGlyLeuLeuGlyProGluTyrrValLysLeuLeuGlyLeu 28

DB 115 GACAGATCTCTCGCTTAAAGGGATCTCGGGCCACAAATATGTAATCTCTGGGCTC 174

QY 29 CysValCysLeuSerGlyCysSer 36

DB 175 TGTGTGTGACTGAGTGGCTGCTCT 198

QY 15

LOCUS AG151043 659 bp DNA linear GSS 09-JAN-2002

DEFINITION Pan troglodytes DNA, clone: RP43-014A14.T7, genomic survey

ACCESSION AG151043

VERSION AG151043.1 GI:16680721

KEYWORDS Pan troglodytes male lymphocytes DNA, clone_11b:RPCT-43 Chimpanzee

SOURCE Male BAC Library clone:RP43-014A14.T7.

ORGANISM Pan troglodytes

REFERENCE 1 Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,

AUTHORS Tokoki,Y., Watanabe,H. and Sakaki,Y.

TITLE BAC end sequences of library RPCT-43

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 659)

AUTHORS Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,

TITLE Tokoki,Y., Watanabe,H. and Sakaki,Y.

JOURNAL Direct Submission

COMMENT Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical

and Chemical Research (RIKEN), Genomic Sciences Center (GSC);

1-7-22 Suehiro-chou,Tsukumi-Ku, Yokohama, Kanagawa 230-0045, Japan

(E-mail:chimbases@sc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/;

Tel:81-45-503-9111, Fax:81-45-503-9170)

Clones are derived from the chimpanzee BAC library RPCT-43 This BAC

end was generated during the R&D process and may have higher chance

of clone tracking errors.

PRIMERS

Sequencing: T7

LIBRARY

Vector 1 : pBACe3.6

R.Site 1 : EcoRI

R.Site 2 : EcoRI.

Location/Qualifiers

1..659

/organism="Pan troglodytes"

/db_xref="taxon:9598"

/clone="RP43-014A14.T7"

/sex="male"

/cell_type="lymphocytes"

/clone_11b="RPCT-43 Chimpanzee Male BAC Library"

BASE COUNT 129 a 159 c 154 g 217 t

ORIGIN

Alignment Scores:

Pred. No.: 0.00161 Length: 659

Score: 100.00 Matches: 19

Percent Similarity: 82.14% Conservative: 4

Best Local Similarity: 67.86% Mismatches: 5

Query Match: 51.81% Indels: 0

DB: 17 Gaps: 0

US-09-513-999c_COPY_1_37 (1-37) x AG151043 (1-659)

QY 9 AspserPheSerLeuGlnGlyLeuLeuGlyProGluTyrrValLysLeuLeuGlyLeu 28

DB 116 GACAGATCTCTCGCTTAAAGGGATCTCGGGCCACAAATATGTAATCTCTGGGCTC 175

QY 29 CysValCysLeuSerGlyCysSer 36

DB 176 TGTGTGTGACTGAGTGGCTGCTCT 199

Search completed: April 24, 2003, 23:19:32

Job time : 1058 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: April 24, 2003, 22:35:13 ; Search time 42 Seconds
(without alignments)
270.168 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSPALODPSFSLQGLGPEYKVLGICVCLSGCST 37

Scoring table: BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

atched: 441362 segs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame.p2n.model -DEV=xlh
-O=/cg2n2_1/uspro_spool/US09513999/runat_18042003_170937_28379/app_query.fasta.1.199
-DB=Issued_Patents.NA -QFMT=fastcap -SUFFIX=p2n.rni -MINMATCH=0.1 -LOOPL=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blomsum62 -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn1.1.32.ernat.18042003_170937_28379 -NCPU=6 -ICPU=3
-NO_XLPXY -NO_MAP -LARGEQUERY -NEG_SCORES=0 -WAIT -LONGLOG -DEV_TIMECUT=120
-WARN_TIMECUT=30 -THREDS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-Ygapop=10 -Ygapext=0.5 -Delop=6 -Delext=7

Database : Issued_Patents.NA.*

- 1: /cg2n2_6/ptodata/1/ina/5A.COMB.seq.*
- 2: /cg2n2_6/ptodata/1/ina/5B.COMB.seq.*
- 3: /cg2n2_6/ptodata/1/ina/6A.COMB.seq.*
- 4: /cg2n2_6/ptodata/1/ina/6B.COMB.seq.*
- 5: /cg2n2_6/ptodata/1/ina/PCITUS.COMB.seq.*
- 6: /cg2n2_6/ptodata/1/ina/Backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	58	30.1	2085	2	US-08-677-049-1
2	57	29.5	7379	4	US-09-341-587-5
3	56	29.0	2073	4	US-09-033-523-6
4	56	29.0	3120	1	US-08-456-647B-19
5	56	29.0	3120	2	US-08-456-647B-19
6	54.5	28.2	2064	3	US-08-875-944B-1
7	54.5	28.2	2064	4	US-09-116-049-3
8	54.5	28.2	2410	2	US-08-780-835B-1
9	54.5	28.2	2410	4	US-09-303-268-1
10	54.5	28.2	2410	4	US-09-116-049-1
11	54	28.0	16998	4	US-09-676-610B-24
12	53.5	27.7	1308	4	US-09-526-993-4

13	53.5	27.7	2167	4	US-09-526-993-3	Sequence 3, Appli
14	53.5	27.7	2176	1	US-07-778-890A-2	Sequence 2, Appli
15	53.5	27.7	3224	4	US-09-526-993-2	Sequence 2, Appli
16	53.5	27.7	6070	4	US-09-526-993-1	Sequence 1, Appli
17	53.5	27.7	6157	4	US-09-526-993-10	Sequence 10, Appli
18	53.5	27.7	6202	4	US-09-526-993-8	Sequence 8, Appli
19	52.5	27.2	59065	4	US-09-813-817-3	Sequence 3, Appli
20	52.5	27.2	59065	4	US-09-813-817-3	Sequence 3, Appli
21	52	26.9	2055	3	US-08-872-855-1	Sequence 3, Appli
22	52	26.9	2800	3	US-08-872-855-1	Sequence 1, Appli
23	51.5	26.7	1019	4	US-09-177-650-128	Sequence 128, App
24	51.5	26.7	1191	2	US-09-061-337-11	Sequence 11, Appli
25	51.5	26.7	1191	2	US-09-122-129-11	Sequence 11, Appli
26	51.5	26.7	1191	3	US-09-340-991-11	Sequence 11, Appli
27	51.5	26.7	1191	3	US-08-974-609-11	Sequence 11, Appli
28	51.5	26.7	1191	4	US-08-549-098-11	Sequence 11, Appli
29	51.5	26.7	1297	3	US-09-083-521-4	Sequence 4, Appli
30	51.5	26.7	3523	3	US-08-749-527-1	Sequence 4, Appli
31	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appli
32	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appli
33	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appli
34	51	26.4	3390	1	US-08-452-802-26	Sequence 26, Appli
35	51	26.4	3390	1	US-08-452-802-26	Sequence 26, Appli
36	51	26.4	3416	1	US-08-453-742-24	Sequence 24, Appli
37	51	26.4	3416	1	US-08-454-464-24	Sequence 24, Appli
38	51	26.4	3416	1	US-08-453-222-24	Sequence 24, Appli
39	51	26.4	3416	1	US-08-452-802-24	Sequence 24, Appli
40	51	26.4	3588	1	US-07-792-885A-2	Sequence 2, Appli
41	51	26.4	9370	1	US-08-320-559-27	Sequence 27, Appli
42	51	26.4	9370	5	PCT-US94-04496-27	Sequence 27, Appli
43	51	26.4	9391	1	US-08-320-559-25	Sequence 25, Appli
44	51	26.4	9391	3	US-08-545-860D-25	Sequence 25, Appli
45	51	26.4	9391	5	PCT-US94-04496-25	Sequence 25, Appli

ALIGNMENTS

RESULT 1
US-08-677-049-1
Sequence 1, Application US/08677049
Patent No. 5858707
GENERAL INFORMATION:
APPLICANT: Guimaraes, M. Jorge
APPLICANT: Bazan, J. Fernando
APPLICANT: McClanahan, Terrilli K.
TITLE OF INVENTION: PURIFIED MAMMALIAN NUCLEOBASE PERMEASES;
TITLE OF INVENTION: NUCLEIC ACIDS; ANTIBODIES
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESS: DNAX Research Institute
STREET: 901 California Avenue
CITY: Palo Alto
STATE: California
COUNTRY: USA
ZIP: 94304-1104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/677,049
FILING DATE: 03-JUL-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/000,788
FILING DATE: 03-JUL-1996
ATTORNEY/AGENT INFORMATION:
NAME: Ching, Edwin P.
REGISTRATION NUMBER: 34,090
REFERENCE/DOCKET NUMBER: DX0511
TELECOMMUNICATION INFORMATION:

QY 2 GlyGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGlu 21
||| |||||:|||||
Db 432 GGTCATCTTTCTTCATTCATGATGGAACCGGAGCTTGTCCGGATCATTTGACCGCAC 491
QY 22 TyrValLysLeuLeuGlyLeuGlyCysVal 30
||| :||| |||
Db 492 CAAGCTCTGTGTGGAAGACTAACCGTG 518

RESULT 4
US-08-456-647B-19
; Sequence 19, Application US/08456647B
; Patent No. 5811516
; GENERAL INFORMATION:
; APPLICANT: Lemke Ph.D. et al., Greg E.
; TITLE OF INVENTION: PROTEIN-TYROSINE KINASE GENES
; NUMBER OF SEQUENCES: 54
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: US
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/456,647B
; FILING DATE: 02-JUN-1995
; CLASSIFICATION: 530
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 08/237,401
; FILING DATE: 02-MAY-1994
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 07/884,486
; FILING DATE: 15-MAY-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Wetherell Ph.D., John R.
; REGISTRATION NUMBER: 31,678
; REFERENCE/DOCKET NUMBER: 07251/007002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 678-5070
; TELEFAX: (619) 678-5099
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3120 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; IMMEDIATE SOURCE:
; CLONE: Tyro-10
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 485..3047
US-08-456-647B-19

Alignment Scores:
Pred. No.: 66.6 Length: 3120
Score: 56.00 Matches: 12
Percent Similarity: 54.84% Conservative: 5
Best Local Similarity: 38.71% Mismatches: 6
Query Match: 29.02% Indels: 8
DB: 1 Gaps: 1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-08-456-647B-19 (1-3120)
QY 3 GlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGluTyr 22
||| |||||:||||| |||||
Db 91 GGAGCTGTGGCTCTTCAGACTCA-----GGACCGAGGCGAG 126

QY 23 ValLysLeuLeuGlyLeuGlyCysValCysLeuSer 33
||| :||| :||| :|||
Db 127 ATTCATGTTTGTGGGCTGTGATTTGTGTGAC 159

RESULT 5
US-08-237-401A-19
; Sequence 19, Application US/08237401A
; Patent No. 5837448
; GENERAL INFORMATION:
; APPLICANT: Lemke Ph.D. et al., Greg E.
; TITLE OF INVENTION: PROTEIN-TYROSINE KINASE GENES
; NUMBER OF SEQUENCES: 54
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: US
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/237,401A
; FILING DATE: 02-MAY-1994
; CLASSIFICATION: 435
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 07/884,486
; FILING DATE: 15-MAY-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Halle Ph.D., Lisa A.
; REGISTRATION NUMBER: 38,347
; REFERENCE/DOCKET NUMBER: 07251/007001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 678-5070
; TELEFAX: (619) 678-5099
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3120 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; IMMEDIATE SOURCE:
; CLONE: Tyro-10
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 485..3047
US-08-237-401A-19

Alignment Scores:
Pred. No.: 66.6 Length: 3120
Score: 56.00 Matches: 12
Percent Similarity: 54.84% Conservative: 5
Best Local Similarity: 38.71% Mismatches: 6
Query Match: 29.02% Indels: 8
DB: 2 Gaps: 1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-08-237-401A-19 (1-3120)
QY 3 GlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGluTyr 22
||| |||||:||||| |||||
Db 91 GGAGCTGTGGCTCTTCAGACTCA-----GGACCGAGGCGAG 126

QY 23 ValLysLeuLeuGlyLeuGlyCysValCysLeuSer 33
||| :||| :||| :|||
Db 127 ATTCATGTTTGTGGGCTGTGATTTGTGTGAC 159

RESULT 6
US-08-875-944B-1/c
; Sequence 1, Application US/08875944B

```
Patent No. 6096542
GENERAL INFORMATION:
APPLICANT: FUJINAGA, Kei
APPLICANT: YOSHIDA, Koichi
APPLICANT: HIGASHINO, Fumihito
TITLE OF INVENTION: CANCER CONTROL
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK, P.L.L.C.
STREET: 624 Ninth Street N.W., Ste. 300
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20001
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/875,944B
FILING DATE: 07-AUG-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP 07-020173
FILING DATE: 08-FEB-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/JP96/00016
FILING DATE: 09-JAN-1996
ATTORNEY/AGENT INFORMATION:
NAME: BROWDY, Roger L.
REGISTRATION NUMBER: 25,618
REFERENCE/DOCKET NUMBER: FUJINAGA-1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 628-5197
TELEFAX: (202) 737-3528
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2064 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1386
US-08-875-944B-1
Alignment Scores:
Seq. No.: 66 Length: 2064
Score: 54.50 Matches: 15
Percent Similarity: 44.44% Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match: 28.24% Indels: 13
DB: 3 Gaps: 1
US-09-513-999c-7869_copy_1_37 (1-37) x US-08-875-944B-1 (1-2064)
QY 3 GlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGluTyr 22
DB 278 GCGGCTTCCTCTCGACAGACGCGCGGCTCTGCGGGGACTCTGGGCTCTTCTTG 219
QY 23 ValIysLeuLeuGlyLeu-----Cys 29
DB 218 ATCCGTGGTGGTGGGCTGTGGAAAGCTAGATTCTGATGAATGGAATCAAGAACTGC 159
QY 30 ValCysLeuSerGly 34
DB 158 TCATCACTGTCTGCT 144
RESULT 7
US-09-116-049-3/c
Sequence 3, Application US/09116049A
```

```
Patent No. 6248351
GENERAL INFORMATION:
APPLICANT: Hung, Men-Chie
TITLE OF INVENTION: HUMAN PEA3 IS A TUMOR SUPPRESSOR FOR CANCER CELLS
FILE REFERENCE: UTSC:582
CURRENT APPLICATION NUMBER: US/09/116,049A
CURRENT FILING DATE: 1998-07-15
NUMBER OF SEQ ID NOS: 11
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 3
LENGTH: 2064
TYPE: DNA
ORGANISM: Homo sapiens
US-09-116-049-3
Alignment Scores:
Pred. No.: 66 Length: 2064
Score: 54.50 Matches: 15
Percent Similarity: 44.44% Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match: 28.24% Indels: 13
DB: 4 Gaps: 1
US-09-513-999c-7869_copy_1_37 (1-37) x US-09-116-049-3 (1-2064)
QY 3 GlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyProGluTyr 22
DB 278 GCGGCTTCCTCTCGACAGACGCGCGGCTCTGCGGGGACTCTGGGCTCTTCTTG 219
QY 23 ValIysLeuLeuGlyLeu-----Cys 29
DB 218 ATCCGTGGTGGTGGGCTGTGGAAAGCTAGATTCTGATGAATGGAATCAAGAACTGC 159
QY 30 ValCysLeuSerGly 34
DB 158 TCATCACTGTCTGCT 144
RESULT 8
US-08-780-835B-1/c
Sequence 1, Application US/08780835B
Patent No. 5922688
GENERAL INFORMATION:
APPLICANT: Hung, Men-Chie
APPLICANT: Xing, Xiangming
TITLE OF INVENTION: PEA3 is a Tumor Suppressor
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: ARNOLD, WHITE AND DURKEE
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: USA
ZIP: 77210-4433
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/780,835B
FILING DATE: 10-JAN-1997
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Wilson, Mark B.
REGISTRATION NUMBER: 37,259
REFERENCE/DOCKET NUMBER: UTSC500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2410 base pairs
TYPE: nucleic acid
```



```

STRANDEDNESS: single
TOPOLOGY: linear
US-08-780-835B-1

Alignment Scores:
Pred. No.:      80.6          Length:    2410
Score:         54.50        Matches:     15
Percent Similarity: 44.44%   Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match:    28.24%       Indels:    13
DB:             2           Gaps:      1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-08-780-835B-1 (1-2410)
Oy      3 gysrerphealnlghlaspserpheserleunglnlgyleuenglpyproglutyr 22
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      694 ggTgccttccTcctgacagacaggcgggGgtCTGTGGCGGACtCTGgTTCCttCTTG 6355
Oy      23 vallyslseuenglgleu-----Cys 29
:: : :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::
634 ATCTGCTGGTGGGGGCTATGGAAGCTAATTCTTCTGAAATGGAATATAGGACAAACTGC 575
Oy      30 vAlCysleusErgly 34
Db      574 TCATCATCTGTCCGGT 560

RESULT 9
US-09-303-268-1/c
; Sequence 1, Application US/09303268
; Patent No. 6172212
GENERAL INFORMATION:
APPLICANT: Hung, Mien-Chie
Xing, Xiangming
TITLE OF INVENTION: PEA3 is a Tumor Suppressor
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESSES:
ADDRESSEE: ARNOLD, WHITE AND DURKEE
STREET: P.O. Box 4433
City: Houston
State: Texas
Country: USA
ZIP: 77210-4433
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/303,268
FILING DATE: 30-Apr-1999
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/780,835
FILING DATE: 10-JAN-1997
ATTORNEY/AGENT INFORMATION:
NAME: Wilson, Mark B.
REGISTRATION NUMBER: 37,259
REFERENCE/DOCKET NUMBER: UTSC500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2410 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-303-268-1
SEQUENCE DESCRIPTION: SEQ ID NO: 1:

Alignment Scores:
Pred. No.:      80.6          Length:    2410
Score:         54.50        Matches:     15
Percent Similarity: 44.44%   Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match:    28.24%       Indels:    13
DB:             2           Gaps:      1

```

```

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-303-268-1 (1-2410)
Qy      3  GlycerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlnGlyProGluTyr  22
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      694  GGTGGCTTCCTGCTGCGACAGACAGGCGGGGCTGTGCGGGGACTCTGGGGTTCCTCTTG  635
Qy      23  ValLysLeuLeuGlnGlyLeu-----Cys  29
      ::: ::::: |||||
Db      634  ATCCGTGGTGGCGGCTATGGAAGCTAAGTTTCTGATGGAATCAGGAACAACACTGC  575
Qy      30  ValCysLeuSerGly  34
      ||||| |||||
Db      574  TCATCATCTGTCCGGT  560

RESULT 10
US-09-116-049-1/c
; Sequence 1, Application US/09116049A
; Patent No. 6248351
; GENERAL INFORMATION:
; APPLICANT: Hung, Men-Chie
; TITLE OF INVENTION: HUMAN PEAK 3 IS A TUMOR SUPPRESSOR FOR CANCER CELLS
; FILE REFERENCE: US05,582
; CURRENT APPLICATION NUMBER: US/09/116,049A
; CURRENT FILING DATE: 1998-07-15
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 2410
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-116-049-1

Alignment Scores:
Pred. No.:      80.6      Length: 2410
Score:          54.50     Matches: 15
Percent Similarity: 44.44% Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match:      28.24%   Indels: 13
Db:               4       Gaps: 1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-116-049-1 (1-2410)
Qy      3  GlycerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlnGlyProGluTyr  22
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      694  GGTGGCTTCCTGCTGCGACAGACAGGCGGGGCTGTGCGGGGACTCTGGGGTTCCTCTTG  635
Qy      23  ValLysLeuLeuGlnGlyLeu-----Cys  29
      ::: ::::: |||||
Db      634  ATCCGTGGTGGCGGCTATGGAAGCTAAGTTTCTGATGGAATCAGGAACAACACTGC  575
Qy      30  ValCysLeuSerGly  34
      ||||| |||||
Db      574  TCATCATCTGTCCGGT  560

RESULT 11
US-09-676-610B-24/c
; Sequence 24, Application US/09676610B
; Patent No. 6444465
; GENERAL INFORMATION:
; APPLICANT: C. Frank Bennett
; APPLICANT: Jacqueline Wyatt
; APPLICANT: Susan M. Freier
; TITLE OF INVENTION: OLIGONUCLEOTIDE INHIBITION OF HER-1 EXPRESSION
; FILE REFERENCE: PUS-0138
; CURRENT APPLICATION NUMBER: US/09/676,610B
; CURRENT FILING DATE: 2000-09-29
; NUMBER OF SEQ ID NOS: 182
; SEQ ID NO 24

```



```

?       TOPOLOGY: circular
?       MOLECULE TYPE: cDNA to genomic RNA
?       HYPOTHETICAL: yes
?       ANTI-SENSE: no
?       ORIGINAL SOURCE:
?       ORGANISM: Newcastle disease virus (NDV)
?       STRAIN: Texas strain
?       INDIVIDUAL ISOLATE: chicken
?       IMMEDIATE SOURCE:
?       LIBRARY: plasmid
?       CLONE: pMDV 108
?       FEATURE:
?       NAME/KEY: part of matrix protein gene and fusion protein gene
?       LOCATION: from to description
?       LOCATION: 1 271 matrix protein gene
?       LOCATION: 431 2092 fusion protein gene
?
US-07-778-890A-2
?
Alignment Scores:
Pred. No.: 100 Length: 2176
Score: 53.50 Matches: 14
Percent Similarity: 55.88 Conservative: 5
Best Local Similarity: 41.18 Mismatches: 12
Query Match: 27.728 Indels: 3
DB: 1 Gaps: 2
?
US-09-513-999C-7869_COPY_1_37 (1-37) x US-07-778-890A-2 (1-2176)
QY 7 LeuGlnpSerpSerSerLeuGlnGly---LeuLeuGlyProGluTyrValysLeu 25
Db 166 TTGATCATGATTTTACACTCCGCGCGCCGACGTTGACTCCAGATATCTTAGCTAC 107
QY 26 Leu-----GlyLeuGlyValysLeuSerGlyCysSerThr 37
Db 106 CTGAGGAGAGCATTCGTCGTATGAGATAGCAGCGCTGCCACT 65
?
RESULT 15
US-09-526-993-2/c
? Sequence 2, Application US/09526993
? Patent No. 6465715
? GENERAL INFORMATION:
? APPLICANT: Zwaal, Richard
? APPLICANT: Aaser, Wouter
? APPLICANT: Roelens, Ingele
? APPLICANT: Bogaert, Thierry
? TITLE OF INVENTION: EXPRESSION OF DNA OR PROTEINS IN C. ELEGANS
? FILE REFERENCE: B0192/7012/ERC/KA
? CURRENT APPLICATION NUMBER: US/09/526,993
? CURRENT FILING DATE: 2000-03-16
? EARLIER APPLICATION NUMBER: U.K. 9906018.8
? EARLIER FILING DATE: 1999-03-16
? NUMBER OF SEQ ID NOS: 11
? SOFTWARE: FastSeq for Windows Version 3.0
? SEQ ID NO 2
? LENGTH: 3224
? TYPE: DNA
? ORGANISM: Caenorhabditis Elegans
US-09-526-993-2
?
Alignment Scores:
Pred. No.: 167 Length: 3224
Score: 53.50 Matches: 11
Percent Similarity: 64.29 Conservative: 1
Best Local Similarity: 39.29 Mismatches: 7
Query Match: 27.728 Indels: 3
DB: 4 Gaps: 1
?
US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-526-993-2 (1-3224)
QY 5 PheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTyrValys 24
Db 1251 TTTCGATTTCAAAACAGATTCGTTTCATATCGAA-----ATTCCGATATTCATTCGC 1201

```

Qy 25 LeuLeuGlyLeuCysValCysLeu 32
||||| : : |||
Db 1200 CTCCTGAAATATTGTCGATGTTG 1177

Search completed: April 24, 2003, 23:20:43
Job time : 58 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: April 24, 2003, 23:01:58 ; Search time 77 Seconds
(without alignments)
522.867 Million cell updates/sec

Title: US-09-513-999C-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSFALQDSSSLQGLGPRYKVLGICVCLSGCST 37

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 709820 seqs, 544064369 residues
Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame_plus_p2n.model -DEV=xlh
-O=/cgn2_1/USPTO_SPOOL/US09513999/runat_18042003_170939_28423/app_query.fasta.1.199
-DB=Published_Applications_NA -OPMT=fastlap -SUFFIX=p2n.rnpb -MINMATCH=0.1
-LOOPEXT=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=bl0sum62
-TRANS=human40.cdi -LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100
-THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEADSIZE=500 -MINLEN=0
-MAXLEN=2000000000 -USER=US09513999 @cgn1.1.1.77 @runat.18042003_170939_28423
-NUPU=6 -ICPR=3 -NO_XLPXY -NO_MAP -LARGEUDERY -NEG_SCORES=0 -WAIT -LONKLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : Published Applications_NA:*

- 1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/ptodata/1/pubpna/PCT_NEW_PUB.seq:*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
- 6: /cgn2_6/ptodata/1/pubpna/PCTUS_PUBCOMB.seq:*
- 7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
- 8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
- 9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
- 10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
- 12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
- 13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
- 14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	93	48.2	684973	10 US-09-263-959-1	Sequence 1, Appli
C 2	88.5	45.9	570	10 US-09-864-761-9118	Sequence 9118, Ap
C 3	78	40.4	428	9 US-09-918-995-8153	Sequence 8153, Ap
C 4	69.5	36.0	412	10 US-09-783-590-1450	Sequence 1450, Ap

5	63	32.6	162	10 US-09-864-761-31393	Sequence 31393, A
6	63	32.6	519	10 US-09-864-761-14866	Sequence 14866, A
7	60.5	31.3	2730	9 US-09-965-528-27	Sequence 27, Appl
8	60.5	31.3	24533	9 US-09-764-868-1349	Sequence 1349, Ap
9	59.5	30.8	506	10 US-09-783-550-5391	Sequence 5391, Ap
C 10	57.5	29.8	1605	10 US-09-930-218-10	Sequence 8, Appl1
11	57	29.5	1155	10 US-09-833-381-8	Sequence 685, App
12	56	29.0	1380	10 US-09-974-300-685	Sequence 19, Appl
13	56	29.0	3120	9 US-09-158-722-19	Sequence 154, App
14	56	29.0	302250	10 US-09-962-832-154	Sequence 11, Appl
C 15	55	28.5	250	10 US-09-044-604-11	Sequence 8266, Ap
C 16	55	28.5	360	10 US-09-867-701-8266	Sequence 4049, Ap
C 17	55	28.5	435	9 US-09-796-692-4049	Sequence 101, App
18	55	28.5	467	10 US-09-864-761-6467	Sequence 1, Appl1
19	55	28.5	747	12 US-10-001-879-101	Sequence 22, Appl
C 20	55	28.5	855	10 US-09-044-604-1	Sequence 541, App
21	55	28.5	3824	9 US-10-036-041-22	Sequence 541, App
22	55	28.5	3824	9 US-10-028-072-541	Sequence 22, Appl
23	55	28.5	3824	9 US-10-035-855-22	Sequence 541, App
24	55	28.5	3824	9 US-10-121-049-541	Sequence 541, App
25	55	28.5	3824	9 US-10-123-904-541	Sequence 541, App
26	55	28.5	3824	9 US-10-140-470-541	Sequence 22, Appl
27	55	28.5	3824	9 US-09-931-836-22	Sequence 541, App
28	55	28.5	3824	9 US-10-175-746-541	Sequence 541, App
29	55	28.5	3824	9 US-10-176-918-541	Sequence 541, App
30	55	28.5	3824	9 US-10-176-921-541	Sequence 541, App
31	55	28.5	3824	9 US-10-227-884-209	Sequence 209, App
32	55	28.5	3824	9 US-10-036-214-22	Sequence 22, Appl
33	55	28.5	3824	9 US-10-137-865-541	Sequence 541, App
34	55	28.5	3824	9 US-10-140-474-541	Sequence 541, App
35	55	28.5	3824	9 US-10-035-719-22	Sequence 22, Appl
36	55	28.5	3824	9 US-10-142-431-541	Sequence 541, App
37	55	28.5	3824	9 US-10-143-114-541	Sequence 541, App
38	55	28.5	3824	9 US-10-230-163-209	Sequence 209, App
39	55	28.5	3824	9 US-10-140-002-541	Sequence 541, App
40	55	28.5	3824	9 US-10-036-160-22	Sequence 22, Appl
41	55	28.5	3824	9 US-10-142-419-541	Sequence 541, App
42	55	28.5	3824	9 US-10-218-631-209	Sequence 209, App
43	55	28.5	3824	9 US-10-230-338-209	Sequence 209, App
44	55	28.5	3824	9 US-10-035-958-22	Sequence 22, Appl
45	55	28.5	3824	9 US-10-036-150-22	Sequence 22, Appl

ALIGNMENTS

RESULT 1
US-09-263-959-1/C
; Sequence 1, Application US/09263959
; Patent No. US20020150891A1
; GENERAL INFORMATION:
; APPLICANT: Hood, Leroy E.
; APPLICANT: Rowen, Lee
; APPLICANT: Koop, Ben F.
; TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U
; NUMBER OF SEQUENCES: 1279
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seed and Berry LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: US
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/263,959
; FILING DATE: 05-MAR-1999
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: McMasters, David D.

```
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 684973 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-263-959-1

Alignment Scores:
Pred. No.: 0.0381 Length: 684973
Score: 93.00 Matches: 19
Percent Similarity: 75.00% Conservative: 2
Best Local Similarity: 67.86% Mismatches: 7
Query Match: 48.15% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-263-959-1 (1-684973)
QY 9 AspSerPheSerSerLeuGlnGlyLeuGlyProGluTyrValIysLeuGlyLeu 28
Db 404731 GACGAGATTCTGCTGCTGCTGGGATCTGAGCGTGGATGTGTAACCTCCTGGGCTCTC 404672
QY 29 CysValCysLeuSerGlyCysSer 36
Db 404671 TGTGTGCTGTGAGCAGCTGCTCT 404648

RESULT 2
US-09-864-761-9118/C
: Sequence 9118, Application US/09864761
: Patent No. US20020048763A1
: GENERAL INFORMATION:
: APPLICANT: Penn, Sharon G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: APPLICANT: Chen, Wensheng
: TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
: FILE REFERENCE: Aecmice-X-1
: CURRENT APPLICATION NUMBER: US/09/864,761
: CURRENT FILING DATE: 2001-05-23
: PRIOR APPLICATION NUMBER: US 60/180,312
: PRIOR FILING DATE: 2000-02-04
: PRIOR APPLICATION NUMBER: US 60/207,456
: PRIOR FILING DATE: 2000-05-26
: PRIOR APPLICATION NUMBER: US 09/632,366
: PRIOR FILING DATE: 2000-08-03
: PRIOR APPLICATION NUMBER: GB 24263.6
: PRIOR FILING DATE: 2000-10-04
: PRIOR APPLICATION NUMBER: US 60/236,359
: PRIOR FILING DATE: 2000-09-27
: PRIOR APPLICATION NUMBER: PCT/US01/00666
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00667
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00664
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00669
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00665
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00668
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00663
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00662
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00661
: PRIOR FILING DATE: 2001-01-30
```

```
PRIOR APPLICATION NUMBER: PCT/US01/00670
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: US 60/234,687
: PRIOR FILING DATE: 2000-09-21
: PRIOR APPLICATION NUMBER: US 09/608,408
: PRIOR FILING DATE: 2000-06-30
: PRIOR APPLICATION NUMBER: US 09/774,203
: PRIOR FILING DATE: 2001-01-29
: NUMBER OF SEQ ID NOS: 49117
: SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
: SEQ ID NO 9118
: LENGTH: 570
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: OTHER INFORMATION: MAP TO AP000053.1
: OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.8
: OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
: OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
: OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
: OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
: OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3
: OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
: OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
US-09-864-761-9118

Alignment Scores:
Pred. No.: 2.61e-05 Length: 570
Score: 88.50 Matches: 21
Percent Similarity: 61.11% Conservative: 1
Best Local Similarity: 58.33% Mismatches: 7
Query Match: 45.85% Indels: 7
Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-864-761-9118 (1-570)
QY 1 MetGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
Db 568 ATGAAATGAGTCT-----CTTCCTTGGGATTCCTTGGGCTG 530
QY 21 GltTyrValIysLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
Db 529 GAGTATGTAAATTCCTGCGCTTGTGTGCTGTGAGCGGCTCT 482

RESULT 3
US-09-918-995-8153
: Sequence 8153, Application US/09918995
: Publication No. US20030073623A1
: GENERAL INFORMATION:
: APPLICANT: Hyseq, Inc.
: TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
: FILE REFERENCE: 20411-756
: CURRENT APPLICATION NUMBER: US/09/918,995
: CURRENT FILING DATE: 2001-07-30
: PRIOR APPLICATION NUMBER: US/09/235,076
: PRIOR FILING DATE: 1999-01-20
: NUMBER OF SEQ ID NOS: 38054
: SOFTWARE: FASTSEQ for Windows Version 3.0
: SEQ ID NO 8153
: LENGTH: 428
: TYPE: DNA
: ORGANISM: Homo sapiens
US-09-918-995-8153

Alignment Scores:
Pred. No.: 0.000899 Length: 428
Score: 78.00 Matches: 17
Percent Similarity: 71.43% Conservative: 3
Best Local Similarity: 60.71% Mismatches: 8
Query Match: 40.41% Indels: 0
Gaps: 0
```

```

1  US-09-513-9996-7869_COPY_1_37 (1-37) x US-09-918-995-8153 (1-428)
2  QY 6 AlaiendinspearheseSesSesLeuGngLeuGngLeuGngProGluTyrValIysLeu 25
3  Db 273 GCCTTGCTGCTATGCCATACCTTTCTTGGGGATCTTAAGGCTGAGTAATGAAGCTC 332
4  QY 26 LeuGlyLeuGlyValCysLeuSer 33
5  Db 333 CTGGGCTCTGTATGTGCTGAAC 356
6
7  RESULT 4
8  US-09-783-590-1450
9  ; Sequence 1450, Application US/09783590
10 ; Patent No. US20020110850A1
11 ; GENERAL INFORMATION:
12 ; APPLICANT: Dillon, Patrick J.
13 ; APPLICANT: Haseltine, William A.
14 ; APPLICANT: Li, Haodong
15 ; APPLICANT: Rosen, Craig A.
16 ; APPLICANT: Ruben, Steven M.
17 ; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
18 ; FILE REFERENCE: PO-16,2C1
19 ; CURRENT APPLICATION NUMBER: US/09/783,590
20 ; CURRENT FILING DATE: 2000-02-15
21 ; PRIOR APPLICATION NUMBER: 08/420,856
22 ; PRIOR FILING DATE: 1995-04-12
23 ; PRIOR APPLICATION NUMBER: 08/246,731
24 ; PRIOR FILING DATE: 1994-11-21
25 ; NUMBER OF SEQ ID NOS: 12485
26 ; SOFTWARE: PatentIn Ver. 2.0
27 ; SEQ ID NO 1450
28 ; LENGTH: 412
29 ; TYPE: DNA
30 ; ORGANISM: Homo sapiens
31 ; FEATURE:
32 ; NAME/KEY: misc feature
33 ; LOCATION: (5)
34 ; OTHER INFORMATION: n equals a,t,g, or c
35 ; NAME/KEY: misc feature
36 ; LOCATION: (189)
37 ; OTHER INFORMATION: n equals a,t,g, or c
38 ; NAME/KEY: misc feature
39 ; LOCATION: (190)
40 ; OTHER INFORMATION: n equals a,t,g, or c
41 ; NAME/KEY: misc feature
42 ; LOCATION: (231)
43 ; OTHER INFORMATION: n equals a,t,g, or c
44 ; NAME/KEY: misc feature
45 ; LOCATION: (265)
46 ; OTHER INFORMATION: n equals a,t,g, or c
47 ; NAME/KEY: misc feature
48 ; LOCATION: (266)
49 ; OTHER INFORMATION: n equals a,t,g, or c
50 ; NAME/KEY: misc feature
51 ; LOCATION: (268)
52 ; OTHER INFORMATION: n equals a,t,g, or c
53 ; NAME/KEY: misc feature
54 ; LOCATION: (269)
55 ; OTHER INFORMATION: n equals a,t,g, or c
56 ; NAME/KEY: misc feature
57 ; LOCATION: (274)
58 ; OTHER INFORMATION: n equals a,t,g, or c
59 ; NAME/KEY: misc feature
60 ; LOCATION: (275)
61 ; OTHER INFORMATION: n equals a,t,g, or c
62 ; NAME/KEY: misc feature
63 ; LOCATION: (339)
64 ; OTHER INFORMATION: n equals a,t,g, or c
65 ; NAME/KEY: misc feature
66 ; LOCATION: (340)
67 ; OTHER INFORMATION: n equals a,t,g, or c
68 ; NAME/KEY: misc feature
69 ; LOCATION: (370)
70 ;

```

```

OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (399)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (402)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (403)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (406)
OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-1450

Alignment Scores:
Pred. NO.:          0.0202           Length:         412
Score:              69.50            Matches:          16
Percent Similarity: 61.29%           Conservative:     3
Best Local Similarity: 51.61%        Mismatches:      7
Query Match:       36.01%            Indels:          5
DB:                10               Gaps:            1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-783-590-1450 (1-412)
OY    11 PheserSerLeuIn-----GlyLeuLeuGlyProGluTyValIysLeu 25
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db    172 TTTCGTGAAGTGCACCTNNNTTTTCTGGGAAGCCCTCGAAAGCCAGAGTATCTGAGGCTN 231
OY    26 LeuGlyLeuCysValCylsLeuSerGlyCysSer 36
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db    232 TTGAATCTGTGTCACAGGCCCTCAGTGCGTCTCT 264

RESULT 5
US-09-864-761-31393
Sequence 31393, Application US/09864761
Patent NO. US2002048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aemica-X-1
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30

```

```

; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
; SEQ ID NO 31393
; LENGTH: 162
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC017089.2
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
; OTHER INFORMATION: SWISSPROT HIT: P98161, EVALUE 3.00e+00
; OTHER INFORMATION: EST_HUMAN HIT: A1792950.1, EVALUE 6.00e-07
; OTHER INFORMATION: NT HIT: AL163210.2, EVALUE 4.00e-04
; US-09-864-761-31393

Alignment Scores:
Pred. No.: 0.0695 Length: 162
Score: 63.00 Matches: 13
Percent Similarity: 66.67% Conservative: 1
Best Local Similarity: 61.90% Mismatches: 7
Query Match: 32.64% Indels: 0
DB: 10 Gaps: 0

Oy 36 Ser 36
|||
Db 120 TCT 122

RESULT 6
US-09-864-761-14866
; Sequence 14866, Application US/09864761
; Patent No. US2002048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aecmica-x-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263,6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
```

```

; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
; SEQ ID NO 14866
; LENGTH: 519
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC017089.2
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
; US-09-864-761-14866

Alignment Scores:
Pred. No.: 0.303 Length: 519
Score: 63.00 Matches: 13
Percent Similarity: 66.67% Conservative: 1
Best Local Similarity: 61.90% Mismatches: 7
Query Match: 32.64% Indels: 0
DB: 10 Gaps: 0

Oy 36 Ser 36
|||
Db 369 TCT 371

RESULT 7
US-09-965-528-27
; Sequence 27, Application US/09965528
; Publication No. US20020187523A1
; GENERAL INFORMATION:
; APPLICANT: INCYTE GENOMICS, INC.
; APPLICANT: TANG, Y. Tom
; APPLICANT: YUE, Henry
; APPLICANT: LAB, Preeti
; APPLICANT: BURFORD, Neil
; APPLICANT: BANDMAN, Olga
; APPLICANT: BAUMZAI, Mariah R.
; APPLICANT: AZIMZAI, Yaida
; APPLICANT: LU, Dying Alpha M.
; APPLICANT: PATTERSON, Chandra
; TITLE OF INVENTION: EXTRACELLULAR SIGNALING MOLECULES
; FILE REFERENCE: PF-0701 USA
; CURRENT APPLICATION NUMBER: US/09/965,528
; CURRENT FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/134,949
; PRIOR FILING DATE: 1999-05-19
; PRIOR APPLICATION NUMBER: 60/144,270
; PRIOR FILING DATE: 1999-07-15
```



```

; PRIOR APPLICATION NUMBER: 60/146,700
; PRIOR FILING DATE: 1999-07-30
; PRIOR APPLICATION NUMBER: 60/157,508
; PRIOR FILING DATE: 1999-10-04
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: PERL Program
; SEQ ID NO 27
; LENGTH: 2730
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: Incyca ID No. US2002018/523A1 1288847CB1
US-09-965-528-27

Alignment Scores:
Pred. No.: 6.24 Length: 2730
Score: 60.50 Matches: 13
Percent Similarity: 59.26% Conservative: 3
Best Local Similarity: 48.15% Mismatches: 10
Query Match: 31.35% Indels: 1
Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-965-528-27 (1-2730)
Qy 10 SerPheSerLeuGlnGlyLeuLeu--GlyProGluTyValysLeuLeuGlyLeu 28
Db 630 ACCTTCCTAGAGGCTCTCAGGGGCTCTTCAGTCCCATCTATGAGAACTAGTGGGTTC 689

Qy 29 CysValCysLeuSerGlyCys 35
Db 690 TCCCTGATGACCAAGGGGTGT 710

RESULT 8
US-09-764-868-1349
; Sequence 1349, Application US/09764868
; Patent No. US2002016871A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT232
; CURRENT APPLICATION NUMBER: US/09/764,868
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - refer to PAM or file wrapper
; NUMBER OF SEQ ID NOS: 1510
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1349
; LENGTH: 24533
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-868-1349

Alignment Scores:
Pred. No.: 100 Length: 24533
Score: 60.50 Matches: 16
Percent Similarity: 60.00% Conservative: 5
Best Local Similarity: 45.71% Mismatches: 13
Query Match: 31.35% Indels: 1
Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-764-868-1349 (1-24533)
Qy 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyPro 20
Db 14224 CTGGGCTGGGCTGTGGCCCTTCTAGCGCTCGTGGTGGCGCCAGAGGCTTTGGGCTCT 14283

Qy 21 GluTyValysLeuLeuGlnGlyLeuGlyCysValCysLeuSerGlyCys 35
Db 14284 GAG--ATGCAACTGCTTGAGCTGAGCCCGGAGATGATAGGCTGC 14325

RESULT 9
US-09-783-590-5991
; Sequence 5991, Application US/09783590

```

```

; Patent No. US20020110850A1
; GENERAL INFORMATION:
; APPLICANT: Dillon, Patrick J.
; APPLICANT: Haseltine, William A.
; APPLICANT: Li, Haodong
; APPLICANT: Rosen, Craig A.
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
; FILE REFERENCE: PO-16,201
; CURRENT APPLICATION NUMBER: US/09/783,590
; CURRENT FILING DATE: 2000-02-15
; PRIOR APPLICATION NUMBER: 08/420,856
; PRIOR FILING DATE: 1995-04-12
; PRIOR APPLICATION NUMBER: 08/346,731
; PRIOR FILING DATE: 1994-11-21
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5991
; LENGTH: 506
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (110)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (331)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (351)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (364)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (402)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (435)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (460)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (502)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-5991

Alignment Scores:
Pred. No.: 1.08 Length: 506
Score: 59.50 Matches: 16
Percent Similarity: 55.26% Conservative: 5
Best Local Similarity: 42.11% Mismatches: 8
Query Match: 30.83% Indels: 9
Gaps: 2

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-783-590-5991 (1-506)
Qy 3 GlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeu----- 18
Db 134 GGAGAAAAAGGAGCAAGATTAAGTTCGCTTCGCGTGGCCCTTCACAGCAAGTGTGG 193

Qy 19 -----GlyProGluTyValysLeuLeuGlnGlyLeuGlyCysValCysLeu 32
Db 194 GGAATTCACTAGAGTGAAGAAATAC---AAATTAATTGGAGGTTCAATCTGTTTG 244

RESULT 10
US-09-930-218-10/c
; Sequence 10, Application US/09930218
; Patent No. US20020034810A1
; GENERAL INFORMATION:
; APPLICANT: goldsmith, orit
; APPLICANT: pecker, iris

```

```
APPLICANT: vlodavsky, israel
APPLICANT: israel, michal
TITLE OF INVENTION: AVIAN AND REPTILE DERIVED POLYNUCLEOTIDE ENCODING A POLYPEPTIDE
TITLE OF INVENTION: HEPARINASE ACTIVITY
FILE REFERENCE: 01/22335
CURRENT APPLICATION NUMBER: US/09/930,218
CURRENT FILING DATE: 2001-08-16
PRIOR APPLICATION NUMBER: 09/666,390
PRIOR FILING DATE: 2000-09-20
NUMBER OF SEQ ID NOS: 16
SOFTWARE: PatentIn version 3.1
SEQ ID NO: 10
LENGTH: 1605
TYPE: DNA
ORGANISM: Gallus gallus
US-09-930-218-10
```

```
Alignment Scores:
Pred. No.: 9.73 Length: 1605
Score: 57.50 Matches: 11
Percent Similarity: 73.68% Conservative: 3
Best Local Similarity: 57.89% Mismatches: 4
Query Match: 29.79% Indels: 1
DB: 10 Gaps: 1
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-930-218-10 (1-1605)

```
OY 20 ProglutryValIysLeuLgIleuGlyLeuGlyCys---ValcysLeuSergIyCysSerThr 37
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 819 CCAGATTTCATGAGAGCTTGAGCAGGTGCTGGCTGGCTGGCTGGCGGGCTGCCACG 763
```

```
RESULT 11
US-09-833-381-8
; Sequence 8, Application US/09833381
; Patent No. US20020132090A1
; GENERAL INFORMATION:
; APPLICANT: Robison, Ketch E.
; FILE OF INVENTION: NO. US20020132090A1el Nucleic Acid and Protein Homologs
; TITLE REFERENCE: 5800-119
; CURRENT APPLICATION NUMBER: US/09/833,381
; CURRENT FILING DATE: 2001-04-11
; PRIOR APPLICATION NUMBER: 09/516,448
; PRIOR FILING DATE: 2000-02-29
; NUMBER OF SEQ ID NOS: 2050
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 1155
; TYPE: DNA
; ORGANISM: Homo sapiens
09-833-381-8
```

```
Alignment Scores:
Pred. No.: 7.73 Length: 1155
Score: 57.00 Matches: 12
Percent Similarity: 53.66% Conservative: 10
Best Local Similarity: 29.27% Mismatches: 13
Query Match: 29.53% Indels: 6
DB: 10 Gaps: 1
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-833-381-8 (1-1155)

```
OY 1 MetcIySerPheAlaLeuGlnAspSerPheSerSerLeu----- 14
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 535 GTAGAGAGCACTATACCTTGATCATGCCAATAATGCGCCTCGCTCGAGCTCTGCAG 594
OY 15 GlnGlyLeuLgIleuGlyProGluTrValIysLeuLgIleuGlyLeuGlyValcysLeuSergIy 34
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 595 CACAGGCTACAGTACCTCGGTACATGAGGCCCTTGCCCTGTGCATCTGTGGCGCTGT 654
```

```
OY 35 Cys 35
|||
DB 655 TGT 657
```

```
RESULT 12
US-09-974-300-685
; Sequence 685, Application US/09974300
; Patent No. US20020146721A1
; GENERAL INFORMATION:
; APPLICANT: Berka, Randy M.
; APPLICANT: Clausen, Id Groth
; TITLE OF INVENTION: Methods For Monitoring Multiple Gene
; TITLE OF INVENTION: Expression
; FILE REFERENCE: 10085,500-US
; CURRENT APPLICATION NUMBER: US/09/974,300
; CURRENT FILING DATE: 2001-10-05
; PRIOR APPLICATION NUMBER: 09/680,598
; PRIOR FILING DATE: 2000-10-06
; PRIOR APPLICATION NUMBER: 60/279,526
; PRIOR FILING DATE: 2001-03-27
; NUMBER OF SEQ ID NOS: 8481
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 685
; LENGTH: 1380
; TYPE: DNA
; ORGANISM: Bacillus licheniformis
US-09-974-300-685
```

```
Alignment Scores:
Pred. No.: 14 Length: 1380
Score: 56.00 Matches: 9
Percent Similarity: 75.00% Conservative: 3
Best Local Similarity: 56.25% Mismatches: 4
Query Match: 29.02% Indels: 0
DB: 10 Gaps: 0
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-974-300-685 (1-1380)

```
OY 20 ProglutryValIysLeuLgIleuGlyLeuGlyValcysLeuSergIyCys 35
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 564 CCGAGCTTGTTCATCTTGCGGAGCTTGCACTGCATCAGCAGATGT 611
```

```
RESULT 13
US-09-158-722-19
; Sequence 19, Application US/09158722
; Publication No. US20030013848A1
; GENERAL INFORMATION:
; APPLICANT: Lemke Ph.D. et al., Greg E.
; TITLE OF INVENTION: PROTEIN-TYROSINE KINASE GENES
; NUMBER OF SEQUENCES: 54
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: US
ZIP: 92037
```

```
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/158,722
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/456,647
FILING DATE: 02-JUN-1995
APPLICATION NUMBER: US 08/237,401
FILING DATE: 02-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/884,486
FILING DATE: 15-MAY-1992
ATTORNEY/AGENT INFORMATION:
NAME: Wetherell Ph.D., John R.
REGISTRATION NUMBER: 31,678
```

```

REFERENCE/DOCKET NUMBER: 07251/007002
TELECOMMUNICATION INFORMATION:
    TELEPHONE: (619) 678-5070
    TELEFAX: (619) 678-5099
    INFORMATION FOR SEQ ID NO: 19:
        SEQUENCE CHARACTERISTICS:
            LENGTH: 3120 base pairs
            TYPE: nucleic acid
            STRANDEDNESS: single
            TOPOLOGY: linear
        MOLECULE TYPE: DNA
        IMMEDIATE SOURCE:
            CLONE: Tyro-10
        FEATURE:
            NAME/KEY: CDS
            LOCATION: 485..3047
US-09-158-722-19

Alignment Scores:
      Seq. No.:          39..4           Length:          3120
      Score:             56..00           Matches:           12
      Percent Similarity: 54..84%         Conservative:     5
      Best Local Similarity: 38..71%       Mismatches:      6
      Query Match:       29.02%            Indels:          8
      DB:                 9               Gaps:           1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-158-722-19 (1-3120)
Oy      3 GlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlnGlyProGluTyr 22
      |||.....|||.....|||.....|||.....|||.....|||.....|||.....|||.....
Db      91 GGGAGCTGGCGCTCTTCACAGACTCA-----GGACCAGAGCGCAG 126

Oy      23 ValLysLeuLeuGlyLeuCysValCysLeuSer 33
      :::::||||:||||:||||:||||:||||:||||:||||:||||:||||:||||:||||:
Db      127 ATCTCATGTTTGGGGCTGGATTGTGTGCAGC 159

RESULT 14
US-09-962-832-154
; Sequence 154: Application US//09962832
; Patent No. US20020110821A1
; GENERAL INFORMATION:
; APPLICANT: Ebnert, Reinhard
; TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using Signature
; FILE REFERENCE: 689290-74
; CURRENT APPLICATION NUMBER: US/09/962.832
; PRIORITY FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/60/235,077
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,280
; PRIOR FILING DATE: 2000-09-25
; NUMBER OF SEQ ID NOS: 259
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 154
; LENGTH: 302250
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-962-832-154

Alignment Scores:
      Pred. No.:          1..27e+04       Length:          302250
      Score:             56..00           Matches:           12
      Percent Similarity: 61..29%         Conservative:     7
      Best Local Similarity: 38..71%       Mismatches:      10
      Query Match:       29.02%            Indels:          2
      DB:                 10              Gaps:           1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-962-832-154 (1-302250)
Oy      7 LeuGlnAspSerPheSerSerLeuGlnGlyLeu-----LeuGlyProGluTyrValLys 24
      |||.....|||.....|||.....|||.....|||.....|||.....|||.....|||.....
Db      174135 CTTGAGGAGTGTGTCGAATCCCTTAATTGCAGCACGCTGTGGGTGGGTGTGTGTG 174194

```

```

Oy      25 LeuleuglYleucysValCyleusSeuGlYcys   35
        :::|::||::|||::|||||::||| |
Db 174195 TGtGTGGGCGTGTGTCTGTCTTTTGtGGGTGT 174227


RESULT    15
US-09-044-604-11/c
Sequence 11, Application us/09044604
Patent No. US20020009718A1
GENERAL INFORMATION:
APPLICANT: TIMMS, Kathy L.
TITLE OF INVENTION: ENDOMETRIOSIS-SPECIFIC SECRETORY
PROTEINS
TITLE OF INVENTION: PROTEINS
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: KOHN & ASSOCIATES
STREET: 30500 NO. US20020009718A1Western Hwy. Suite 410
City: Farmington Hills
STATE: Michigan
COUNTRY: US
ZIP: 48334
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/044,604
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Montgomery, Ilene N.
REGISTRATION NUMBER: 38,972
REFERENCE/DOCKET NUMBER: 0994,00084
TELECOMMUNICATION INFORMATION:
TELEPHONE: (248) 539-5050
TELEFAX: (248) 539-5055
INFORMATION FOR SEQ ID NO.: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 250 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "Probefor human ENDO-I"
HYPOTHETICAL: NO
ANTI-SENSE: NO
US-09-044-604-11

Alignment Scores:
Pred. NO.:          2.35           Length:       250
Score:             55.00         Matches:       12
Percent Similarity: 51.61%     Conservative:  4
Best local Similarity: 38.71%   Mismatches:   15
Query Match:       28.50%      Indels:        0
DB:               10              Gaps:          0
US-09-513-9990-7869_COPY_1_37 (1-37) x US-09-044-604-11 (1-250)
OY      2 GlylyserPhehalalenaLnsperPheserSerLeuclnSlneuDeuClProclu 21
        ||| {}{}:: :::::: ||| {}|{|}|{||}||| |||
Db 191 GGAGTGTAATTCGGGTGTACAAACAACCCTTCAACTCTCAAGAAGTCCCTTTGCCCAT 133
OY      22 TyrvallYSleueuqlYleucysvalCYslieu 32
        ||| {}{} ||| {}{}:: |||
DB 131 AGAGTGTAAGTAGGGAATAATCTTCCTGCCTG 99

Search completed: April 24, 2003, 23:57:24
Job time : 153 secs
```


GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:54:24 ; Search time 938 Seconds

(without alignments)
3443.935 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111

Sequence: 1 atgggtgattcttgcctt.....gacctgagtgctgctact 111

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : GenBankl:*

1: gb-ba:*

2: gb-htg:*

3: gb-in:*

4: gb-om:*

5: gb-ov:*

6: gb-pat:*

7: gb-ph:*

8: gb-pl:*

9: gb-pr:*

10: gb-ro:*

11: gb-sts:*

12: gb-sy:*

13: gb-un:*

14: gb-vl:*

15: em-ba:*

16: em-fun:*

17: em-hum:*

18: em-in:*

19: em-mu:*

20: em-om:*

21: em-or:*

22: em-ov:*

23: em-pat:*

24: em-ph:*

25: em-pl:*

26: em-ro:*

27: em-sts:*

28: em-un:*

29: em-vl:*

30: em-htg-hum:*

31: em-htg-inv:*

32: em-htg-other:*

33: em-htg-mus:*

34: em-htg-pln:*

35: em-htg-rod:*

36: em-htg-mam:*

37: em-htg-vtl:*

38: em-sy:*

39: em-htgo-hum:*

40: em-htgo-mus:*

41: em-htgo-other:*

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	169620	2 AC012674	AC012674 Homo sapi
2	66.8	60.2	199038	2 AC116565	AC116565 Homo sapi
3	65.2	58.7	130636	2 AC092586	AC092586 Homo sapi
4	64.6	58.2	140040	9 AC022029	AC022029 Homo sapi
5	64.6	58.2	147288	9 AL355812	AL355812 Human DNA
6	63.6	57.3	75270	9 AL591048	AL591048 Human DNA
7	63.6	57.3	143372	9 AL137847	AL137847 Human DNA
8	63.6	57.3	153053	9 AL137179	AL137179 Human DNA
9	63.6	57.3	153422	2 AC023408	AC023408 Homo sapi
10	63.4	57.1	86314	9 AC109592	AC109592 Homo sapi
11	63.4	57.1	159475	2 AC021378	AC021378 Homo sapi
12	63	56.8	153783	9 AL157937	AL157937 Human DNA
13	62.4	56.2	123779	30 AC021025	AC021025 Homo sapi
14	62.4	56.2	128118	2 AC076969	AC076969 Homo sapi
15	62.4	56.2	128583	2 AC121249	AC121249 Homo sapi
16	62.4	56.2	131347	2 AC002421	AC002421 Homo sapi
17	62.4	56.2	170623	9 AL391375	AL391375 Human DNA
18	62.4	56.2	178650	9 AC104303	AC104303 Homo sapi
19	61.4	55.3	21104	9 AC110811	AC110811 Homo sapi
20	61.4	55.3	121902	9 AC008386	AC008386 Homo sapi
21	61.4	55.3	163083	9 AC103773	AC103773 Homo sapi
22	61.4	55.3	163231	9 AL162575	AL162575 Human DNA
23	61.4	55.3	164819	9 AC092335	AC092335 Homo sapi
24	61.4	55.3	170736	2 AC013658	AC013658 Homo sapi
25	61.4	55.3	177423	9 AC120036	AC120036 Homo sapi
26	61.4	55.3	184866	9 AC016650	AC016650 Homo sapi
27	61.4	55.3	187847	2 AC023155	AC023155 Homo sapi
28	61.4	55.3	196145	9 AC026241	AC026241 Homo sapi
29	60.8	54.8	38936	9 AL358817	AL358817 Human DNA
30	60.8	54.8	169772	2 AC069538	AC069538 Homo sapi
31	60.4	54.4	32918	2 AC007445	AC007445 Homo sapi
32	60.4	54.4	68137	2 AC079966	AC079966 Homo sapi
33	60.4	54.4	159747	2 AP001019	AP001019 Homo sapi
34	60.4	54.4	163065	9 AC021483	AC021483 Homo sapi
35	60.4	54.4	168062	30 AC022292	AC022292 Homo sapi
36	60.4	54.4	169462	9 AC026826	AC026826 Homo sapi
37	60.4	54.4	186224	2 AC105901	AC105901 Homo sapi
38	60.4	54.4	200333	2 AC073279	AC073279 Homo sapi
39	60	54.1	148290	9 AC107979	AC107979 Homo sapi
40	60	54.1	165649	9 AC103996	AC103996 Homo sapi
41	60	54.1	192826	9 AC090762	AC090762 Homo sapi
42	59.8	53.9	138038	9 AL591435	AL591435 Human DNA
43	59.8	53.9	148352	2 AC025309	AC025309 Homo sapi
44	59.8	53.9	171532	2 AC061993	AC061993 Homo sapi
45	59.8	53.9	173260	9 CNS07ECU	AL442183 Human chr

ALIGNMENTS

RESULT 1

AC012674/c AC012674 169620 bp DNA linear HTG 07-SEP-2000

LOCUS Homo sapiens chromosome 3 clone RP1-458H3, WORKING DRAFT SEQUENCE, 18 unordered pieces.

ACCESSION AC012674

VERSION AC012674.10 GI:9719580

KEYWORDS HTG: HTGS_PHASE1; HTGS_DRAFT.

SOURCE

ORGANISM Homo sapiens.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 169620)

Muzny,D.M., Adams,C., Bailey,M., Barabara,J., Blankenburg,K., Bodola,B., Bouck,J., Bowle,S., Brooks,A., Bunay,C., Bunac,C.,

Pred. No. is the number of results predicted by chance to have a

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Donah-Rashid, N., Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D., Forcum-Tansey, J., Frantz, P., Ganesh, R., Gorrell, J.H., Gorrell, L.L., Guenara, M., Harris, K., Hernandez, J., Hodgson, A., Hognes, M., Holloway, C., Hosak, H., Jackson, L.E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondajewski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Lichtarge, O., Liu, J., Liu, W., Logan, O., Lozano, R.J., Lu, J., Lucier, R., Martin, R., Martinez, C., McLeod, M.P., Mei, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Ouellet, C., Parish, B., Paxton, S., Payton, B., Perez, L., Pu, L.L., Sui, E., Shen, H., Simon, M., Sammel, S., Say, J., Scherer, S., Taber, P., Taylor, T., Vasquez, L., Vinson, R., Vo, Q., Webb, M., Watlington, S., Weinstein, G., Weinstein, I.R., Williamson, A., Worley, K., Wren, J., Wrenford, G., Yu, W., Zhou, X., Nelson, D. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.

Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HMOG
Center clone name: RPI-458H3

----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-coverage estimation
Estimated insert size: 171608; agarose-fp estimation
Quality coverage: 3.9x in Q20 bases; agarose-fp estimation
Quality coverage: 4.1x in Q20 bases; sum-of-coverage estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank/draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 28689: contig of 28689 bp in length
28690 28789: gap of unknown length
28790 50832: contig of 22043 bp in length
50833 50932: gap of unknown length
50933 69144: contig of 18212 bp in length
69145 69244: gap of unknown length
69245 84204: contig of 14960 bp in length
84205 84304: gap of unknown length
84305 94667: gap of 10363 bp in length
94668 107261: contig of 12494 bp in length
107262 107361: gap of unknown length
107362 117550: contig of 10189 bp in length
117551 117650: gap of unknown length
117651 126939: contig of 9289 bp in length
126940 127039: gap of unknown length
127040 135040: contig of 8001 bp in length
135041 135140: gap of unknown length
135141 141639: contig of 6499 bp in length

141640 141739: gap of unknown length
141740 149558: contig of 7819 bp in length
149559 149658: gap of unknown length
149659 154562: contig of 4904 bp in length
154563 154662: gap of unknown length
154663 158987: contig of 4325 bp in length
158988 159087: gap of unknown length
159088 162376: contig of 3289 bp in length
162377 162476: gap of unknown length
162477 165191: contig of 2715 bp in length
165192 165291: gap of unknown length
165292 167173: contig of 1882 bp in length
167174 167273: gap of unknown length
167274 168393: contig of 1120 bp in length
168394 168493: gap of unknown length
168494 169620: contig of 1127 bp in length.

Location/Qualifiers
1. 169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RPI-458H3"

BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others

ORIGIN

Query Match 100.0%; Score 111; DB 2; Length 169620;
Best Local Similarity 100.0%; Pred. No. 6.5e-27;
Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGGGTGATCTTTTGGCTTGCAGAGATCTTTTCATCTTTGAGGAGCTTCGGGCGG 60
|||||
Db 87441 ATGGGTGATCTTTTGGCTTGCAGAGATCTTTTCATCTTTGAGGAGCTTCGGGCGG 87382
|||||

OY 61 GAGATGTAACCTCCGGCTCTGTGTGCTGAGTGGCTCTACT 111
|||||
Db 87381 GAGATGTAACCTCCGGCTCTGTGTGCTGAGTGGCTCTACT 87331
|||||

RESULT 2
AC116565/c 199038 bp DNA 1linear HTG 25-APR-2002
AC116565/ Homo sapiens chromosome 4 clone RPI1-1263C18, WORKING DRAFT
AC116565 SEQUENCE, 6 unordered pieces.

ACCESSION AC116565.3 GI:20304073
VERSION AC116565.3
KEYWORDS HTG: HTGS.PHASE1; HTGS.DRAFT; HTGS.ACTIVEFIN.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 199038)
Waterston, R.H.
The sequence of Homo sapiens clone

JOURNAL Unpublished
AUTHORS 2 (bases 1 to 199038)
TITLE Waterston, R.H.
JOURNAL Direct Submission
Submitted (29-MAR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
3 (bases 1 to 199038)
Waterston, R.H.
Direct Submission
Submitted (25-APR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
On Apr 25, 2002 this sequence version replaced gi:20128757.

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc/index.shtml>
Contact: submissions@wustl.wustl.edu

```

----- Project Information -----
Center project name: H_NH1263C18
----- Summary Statistics -----
Sequencing vector: M13, 0%
Sequencing vector: plasmid, 100%
Chemistry: Dye-Primer ET; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 196905 bases at least Q40
Consensus quality: 197086 bases at least Q30
Consensus quality: 197234 bases at least Q20
Insert size: 202000; agarose-ftp
Insert size: 196637; sum-of-ctrls
Quality coverage: 10.73 in Q20 bases; sum-of-ctrls
Quality coverage: 10.82 in Q20 bases; sum-of-ctrls
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 3283: contig of 3283 bp in length
* 3284 3383: gap of unknown length
* 3384 12884: contig of 9501 bp in length
* 12885 12984: gap of unknown length
* 12985 29821: contig of 16837 bp in length
* 29822 29922: gap of unknown length
* 29923 74050: contig of 44129 bp in length
* 74051 74151: gap of unknown length
* 74152 197985: contig of 123835 bp in length
* 197986 198086: gap of unknown length
* 198087 199038: contig of 953 bp in length.
* 199039
Location/Qualifiers
source
1. 199038
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="4"
/clone="RP11-1263C18"
misc_feature
1. 3283
/note="assembly_name:Contig18"
3384. 12884
/note="assembly_name:Contig19"
misc_feature
12985. 29821
/note="assembly_name:Contig20"
29922. 74050
/note="assembly_name:Contig21"
74151. 197985
/note="assembly_name:Contig22"
198086. 199038
/note="assembly_name:Contig14"
misc_feature
199039
BASE COUNT 53552 a 46319 c 46377 g 52290 t 500 others
ORIGIN
Query Match 60.2%; Score 66.8; DB 2; Length 199038;
Best Local Similarity 81.9%; Pred. NO. 7.1e-12;
Matches 77; Conservative 0; Mismatches 17; Indels 0; Gaps 0;
QY 18 CTTGAGAGATTTCTTTTATCTTTTCAGGACCTTGTGGGCGGAGATGATGAATAACCTCT 77
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 58755 CTGGAAGATGATTTCCACCTTCCACGAGATCTGGGGCGGAGATATGTAACGCT 58696
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 78 GGGTCTGTGTGTGTCGAGAGGCTGCTACT 111
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 58695 GGGTCTGTGTGTGTCGAGAGGCTGCTACT 58662
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
RESULT 3
AC092586 AC092586 130636 bp DNA linear PRT 01-MAR-2002
LOCUS

```

```

DEFINITION Homo sapiens BAC clone RP11-30H15 from 2, complete sequence.
ACCESSION AC092586 AC016389
VERSION AC092586.2 GI:15638723
KEYWORDS
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 130636)
AUTHORS Sulston, J.E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
PUBMED 9847074
REFERENCE
2 (bases 1 to 130636)
AUTHORS Haglund, K., Kozlowicz, A., Elliott, G. and Boyer, E.
TITLE The sequence of Homo sapiens BAC clone RP11-30H15
JOURNAL Unpublished (2001)
REFERENCE
3 (bases 1 to 130636)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (19-JUL-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE
4 (bases 1 to 130636)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (18-SEP-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE
5 (bases 1 to 130636)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (01-MAR-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
on Sep 18, 2001 this sequence version replaced gi:14916171.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@wustl.wustl.edu
----- Summary Statistics
Center project name: H_NH030H15
Drafting Center: WIBR
-----

```

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RPI1-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E., Taleno, M., Calanese, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>


```

KEYWORDS      HTG.
SOURCE        Homo sapiens.
ORGANISM      Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE     1 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Genome Therapeutics Corporation Sequencing Center: Human Genome
              Sequence Data
JOURNAL       Unpublished
REFERENCE     2 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (25-JUN-2000) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     3 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (14-JUL-2001) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     4 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (28-MAR-2002) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     5 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (09-APR-2002) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
COMMENT       On Mar 26, 2002 this sequence version replaced gi:14718295.

-----
Center: Genome Therapeutics Corporation
Center code: GTC
Web site: http://www.genomecorp.com/
Contact: gtc-seqcenter@genomecorp.com
Project Information
Center project name: hg107

-----
IMPORTANT: This sequence is the entire
            insert of clone RP11-93L14.
            The true right end of clone RP11-316G7 is at
            2000 in this sequence.

----- Summary Statistics -----
Sequencing vector: N/A
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 990315

-----
Location/Qualifiers
1..140040
   /organism="Homo sapiens"
   /db_xref="taxon:9606"
   /chromosome="10"
   /clone="RP11-93L14"
   /clone_lib="RPCI-11"
BASE COUNT      48331 a 26787 c 24761 g 40161 t
ORIGIN
Query Match          58.2%; Score 64.6; DB 9; Length 140040;
Best Local Similarity 73.9%; Pred. No. 4e-11;
Matches 82; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
Oy      1 ATGGGTGCATCTTTTACCTTGACAGATTCTTTTCATCTTTGACAGAGACTTGTGGGCCG 60
      ||| |||| | |||| | ||| | || |||| || | |||| |||
Db 94202 ATGTAATGACCTCCCGCCTTGACTGACGTAGTAGTACCCTTGCCAGGAGATCCTGTGGCTG 94143
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Oy      61 CAGTATGTAAACATCTCTGGGTCTCTGTGTGTGCTCAATGAGTGGCTGCTACT 111
      | ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 94142 GGATATGAATACTCTCTGGGTCTCTGTGTCAATCACTGAGTGGCTGCTGCT 94092

```

LOCUS	AL355812	147288 bp	DNA	linear	PRI 13-JUN-2001
DEFINITION	Human DNA sequence from clone RP11-81003 on chromosome Xq22.3-24, complete sequence.				
ACCESSION	AL355812				
VERSION	AL355812.23	GI:14456206			
KEYWORDS	HTG.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
AUTHORS	1 (bases 1 to 147288)				
TITLE	Smith,M.				
JOURNAL	Direct Submission				
COMMENT	Submitted (13-JUN-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Requests: clonerequests@sanger.ac.uk On Jun 14, 2001 this sequence version replaced gi:14280423. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TrEMBL; Wp:, WormPep; information on the WormPep database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/ChrX RP11-81003 is from the library RPci-11.3 constructed by the group of Pieter de Jong. For further details see http://www.chori.org/bacpac/home.htm VECTOR: pBAC3.6 IMPORTANT: This sequence is not the entire insert of clone RP11-81003 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone RP11-81003 is at 1 in this sequence. The true left end of clone RP6-200F4 is at 147189 in this sequence.				
FEATURES	Location/Qualifiers				
source	1..147288				
	/organism="Homo sapiens"				
	/db_xref="taxon:9606"				
	/chromosome="X"				
	/map="q22.3-24"				
	/clone="RP11-81003"				
	/clone_lib="RPci-11.3"				
	1381..1674				
repeat_region	/note="Aluub repeat: matches 1..296 of consensus"				
	1757..1937				
repeat_region	/note="Aluo repeat: matches 122..300 of consensus"				
	2002..2645				
repeat_region	/note="L1M4 repeat: matches 2110..2837 of consensus"				
	2649..3323				
repeat_region	/note="L1MC4 repeat: matches 6404..7210 of consensus"				
	3811..3878				
repeat_region	/note="L1MC repeat: matches 2099..2180 of consensus"				
	3882..4044				
repeat_region	/note="L1MC3A repeat: matches 5665..5828 of consensus"				
	4236..4533				
repeat_region	/note="L1MC3A repeat: matches 5257..5563 of consensus"				
	4724..5174				
repeat_region	/note="L1M4 repeat: matches 4562..5026 of consensus"				
	5401..5663				

```
/note="Charliela repeat: matches 1183. .1449 of consensus"
repeat_region 5675. .5805
/note="L1P3 repeat: matches 6014. .6145 of consensus"
repeat_region 5807. .6234
/note="L1 repeat: matches 4970. .5403 of consensus"
repeat_region 6251. .6998
/note="Charliela repeat: matches 404. .1196 of consensus"
repeat_region 6999. .7036
/note="19 copies 2 mer tt 100% conserved"
repeat_region 7244. .7864
/note="L1P repeat: matches 4160. .4781 of consensus"
repeat_region 7865. .8274
/note="Charliela repeat: matches 6. .417 of consensus"
repeat_region 8595. .8673
/note="MER58 repeat: matches 1. .83 of consensus"
repeat_region 8689. .8885
/note="L1M4 repeat: matches 5270. .5464 of consensus"
repeat_region 8942. .11399
/note="L1M4 repeat: matches 2661. .5149 of consensus"
repeat_region 11442. .11832
/note="L1M4 repeat: matches 2153. .2568 of consensus"
repeat_region 11837. .11904
/note="34 copies 2 mer tt 66% conserved"
repeat_region 11917. .12078
/note="L1M2 repeat: matches 5989. .6155 of consensus"
repeat_region 12171. .13851
/note="L1M9 repeat: matches 4533. .6275 of consensus"
repeat_region 13877. .14584
/note="L1M4 repeat: matches 5516. .6298 of consensus"
repeat_region 14582. .16120
/note="L1M1 repeat: matches 1441. .2965 of consensus"
repeat_region 16157. .16267
/note="L1M2 repeat: matches 1666. .1777 of consensus"
repeat_region 16278. .17065
/note="L1M1 repeat: matches 5224. .6053 of consensus"
repeat_region 17066. .17368
/note="Alusx repeat: matches 1. .302 of consensus"
repeat_region 17369. .17599
/note="L1M1 repeat: matches 5010. .5224 of consensus"
repeat_region 17654. .19512
/note="L1M1 repeat: matches 4389. .6262 of consensus"
repeat_region 19745. .20100
/note="L1M2 repeat: matches 5470. .5826 of consensus"
repeat_region 20094. .20104
/note="L1 repeat: matches 4404. .4413 of consensus"
repeat_region 20105. .20666
/note="L1P6 repeat: matches 5582. .6143 of consensus"
repeat_region 20667. .21348
/note="L1 repeat: matches 3734. .4404 of consensus"
repeat_region 21349. .21633
/note="Alusg repeat: matches 1. .291 of consensus"
repeat_region 21634. .22811
/note="L1 repeat: matches 2511. .3734 of consensus"
repeat_region 22853. .23268
/note="L1M4 repeat: matches 3293. .3703 of consensus"
repeat_region 23269. .23420
/note="L1P14 repeat: matches 6004. .6149 of consensus"
repeat_region 23421. .24030
/note="L1M4 repeat: matches 2665. .3293 of consensus"
repeat_region 24021. .24198
/note="L1M4 repeat: matches 2459. .2637 of consensus"
repeat_region 24207. .24312
/note="L1M4 repeat: matches -3. .103 of consensus"
repeat_region 24336. .24651
/note="L1M2C repeat: matches 2291. .2280 of consensus"
repeat_region 24895. .25759
/note="L1M9 repeat: matches 5440. .6304 of consensus"
repeat_region 25760. .26060
/note="Alusx repeat: matches 1. .301 of consensus"
repeat_region 26061. .26155
/note="L1M9 repeat: matches 5346. .5440 of consensus"
repeat_region 26213. .26559
/note="L1M1 repeat: matches 1. .354 of consensus"
```

```
repeat_region 26561. .29502
/note="L1M9 repeat: matches 2349. .5253 of consensus"
repeat_region 29486. .29729
/note="L1M3 repeat: matches 7455. .7734 of consensus"
repeat_region 29737. .30026
/note="L1M4 repeat: matches 2689. .2989 of consensus"
repeat_region 30048. .30693
/note="L1M4 repeat: matches 2187. .2831 of consensus"
repeat_region 30716. .31150
/note="L1P1 repeat: matches 5656. .6155 of consensus"
repeat_region 31299. .31641
/note="L1M7 repeat: matches 5814. .6166 of consensus"
repeat_region 31664. .34500
/note="L1M8 repeat: matches 3262. .6168 of consensus"
repeat_region 34545. .34894
/note="L1M8 repeat: matches 2920. .3279 of consensus"
repeat_region 34901. .35618
/note="L1M8 repeat: matches 5564. .6291 of consensus"
repeat_region 35619. .36010
/note="L1M8 repeat: matches 5204. .5597 of consensus"
repeat_region 36008. .36242
/note="L1M4 repeat: matches 2694. .2931 of consensus"
repeat_region 36243. .36433
/note="L1P13 repeat: matches 5960. .6149 of consensus"
repeat_region 36495. .36721
/note="Tigger3(Golem) repeat: matches 2759. .3038 of consensus"
repeat_region 37064. .37632
/note="L1M4 repeat: matches 2110. .2695 of consensus"
repeat_region 37639. .37766
/note="64 copies 2 mer tt 59% conserved"
repeat_region 38204. .38632
/note="MER5A repeat: matches 1. .445 of consensus"
repeat_region 38696. .38991
/note="L1M4C repeat: matches 1768. .1603 of consensus"
repeat_region 38992. .39287
/note="Alusg repeat: matches 1. .307 of consensus"
repeat_region 39288. .39456
/note="L1M4C repeat: matches 1608. .1769 of consensus"
repeat_region 39501. .40086
/note="L1M4B repeat: matches -273. .334 of consensus"
repeat_region 40087. .40139
/note="L1M7 repeat: matches 5776. .5828 of consensus"
repeat_region 40136. .41315
/note="L1M4 repeat: matches -5. .1210 of consensus"
repeat_region 41348. .42700
/note="L1M4C repeat: matches 940. .2376 of consensus"
repeat_region 42986. .43636
/note="L1M4 repeat: matches 3439. .4077 of consensus"
repeat_region 43654. .44154
/note="L1M2 repeat: matches 4128. .4602 of consensus"
repeat_region 44155. .44465
/note="Alusx repeat: matches 1. .310 of consensus"
repeat_region 44466. .45911
```

Query Match 58.2%; Score 64.6; DB 9; Length 147288;
Best Local Similarity 73.9%; Pred. No. 46-11;
Matches 82; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

```
QY 1 ATGGGTGATCTTTTGCCCTTGACAGATCTTTTCACTTTTGACGAGACTTGGGCGG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 50611 ATGTACGACCTCCGCGCTTCCCTGAGTGCACAGCACTTGTGGGATCTCGGCTCA 50670
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 GAGATGTAACCTCGGCTGCTGAGTGCCTGAGTGCCTGCTACT 111
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 50671 GAGATGTAAGCTCTGGCTCTGTGTGTGCTGAGCAGCTGCTTGTCT 50721
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
```

RESULT 6
AL591048/c AL591048 75270 bp DNA linear PRI 09-AUG-2001
LOCUS Human DNA sequence from clone RP11-612M16 on chromosome 6, complete
DEFINITION sequence.

	ACCESSION	AL591048
	VERSION	AL591048.7
	KEYWORDS	GI:15147695
	SOURCE	Htg.
	ORGANISM	human.
	AUTHORS	Homo sapiens
	TITLE	Euhayrolta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
	JOURNAL	1 (bases 1 to 75270)
	COMMENT	Almeida,J. Direct Submission Submitted (09-AUG-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk Requests: clonerequests@sanger.ac.uk On Aug 10, 2001 this sequence version replaced gi:15020550. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the validation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP. Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/chr6 Rp11-612M16 is from the library RPC1-11.3 constructed by the group of Pieter de Jong. For further details see http://www.chori.org/dacpac/home.htm VECTOR: pBACE3.6 IMPORTANT: This sequence is not the entire insert of clone Rp11-612M16 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone Rp1-206F19 is at 73271 in this sequence. The true right end of clone Rp11-82M9 is at 2000 in this sequence. Location/Qualifiers 1..75270 /organism="Homo sapiens" /db_xref="taxon:9606" /chromosome="6" /clone="Rp11-612M16" /clone_lib="RPC1-11.3" /c1one_lib="RPC1-11.3" 33923..34247 /note="Sequence from reads from a short insert library derived from a single pUC clone. Restriction digest data confirm the assembly." 46760..46849 /note="Sequence from reads from a short insert library derived from a single pUC clone. Restriction digest data confirm the assembly." 49353..49464 /note="Sequence from reads from a short insert library derived from a single pUC clone. Restriction digest data confirm the assembly."
	BASE COUNT	25749 a 14850 c 13893 g 20778 t
	ORIGIN	
	Query Match	57.3%; Score 63.6; DB 9; Length 75270;
	Best Local Similarity	73.6%; Pred. No. 8,7e-11;
	Matches	81; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
1	ATGGGTGAGCTTTTGCGTCAGAGATTCTTTTCATCTTGACGAGCCTTCGGGCCG	60

[illegible]

Db 82342 GATGAGATTTCCTCCCTGGCAGAGATCTGGGGGCGACGATGTGTAACCTCGGGTCTC 82283

QY 85 TGTGTGCTGCTGAGTGGCTGCTCTAC 110

Db 82282 TGTGTGCTGCTGAGTGGCTGCTCTGC 82257

RESULT 8	
LOCUS	AL137179/c
DEFINITION	AL137179 153053 bp DNA linear PRI 30-SEP-2000
ACCESSION	Human DNA sequence from clone RP11-82M9 on chromosome 6, complete sequence.
VERSION	AL137179
KEYWORDS	AL137179..14 GI:10443372
SOURCE	HTG.
ORGANISM	human.
	Homo sapiens
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE	1 (bases 1 to 153053)
AUTHORS	Tracey/A.
TITLE	Direct Submission
JOURNAL	Submitted (26-SEP-2000) Sanger Centre, Hinxton, Cambridgeshire,

COMMENT

corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em1; EMBL; SW: SWISSPROT; Tr1; TREMBL; Wb1; WORMPEP; Information on the WORMPEP database can be found at <http://www.sanger.ac.uk/projects/C.elegans/wormpep> This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger centre Chromosome 6 Mapping group. Further information can be found at

RP11-82m9 is from the library RPI1-11.1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/>

VECTOR: pBACe3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-82m9. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The true right end of clone RP1-82m9 is at 153053 in this sequence. The true right end of clone RP1-27919 is at 100 in this sequence.

FEATURES

```
source
1. 153053
  /organism="Homo sapiens"
  /db_xref="taxon:9606"
  /chromosome="6"
  /clone="RP11-82M9"
  /clone_lib="RPc1-11.1"
102234..102253
```

missing according to restri

ORIGIN

Query Match	57.3%;	Score 63.6;	DB 9;	Length 153053;
Best Local Similarity	73.6%;	Pred. No. 8.7e-11;		

Matches	81: Conservative	0: Mismatches	29: Indels	0: Gaps	0:
Oy	1	ATGGGTGATCTTTTCTCTTGAGGATTCCTTTTATCTTTCAGAGGACTTGTGGGCGG	60		
Db	152945	ATGTCGACAGCTTCTTCCTTCGCTAGACGTAGACATATTTCGTGATATCTGTGGGGCTG	152885		
Oy	61	GAGTATGTAAACTCTGCGGCTCTGTGTGTGTGCGCTGAGTGGCTCTCTTAC	110		
Db	152885	GAGTATGTAAAGCTTCTGTGCTGTGTGTGTGCGCTGACACACTTTTCTGC	152836		

RESULT	9
AC023408	
LOCUS	
DEFINITION	AC023408 153422 bp DNA linear HTG 07-JUN-2006
ACCESSION	Homo sapiens chromosome 6 clone RP11-612M16 map 6, WORKING DRAFT
VERSION	AC023408.2
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE	Homo sapiens.
ORGANISM	Homo sapiens

REFERENCE 1 (bases 1 to 153422)
AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE Homo sapiens chromosome 6, clone RP11-612M6
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 153422)
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.

JOURNAL
REFERENCE
AUTHORS

Unpublished
2 (bases 1 to 153422)

Altren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Bana, N., Beda, F., Boguslavsky, L.,
Boukhalter, B., Brown, A., Burkett, G., Campiano, A., Castle, A.,
Choepel, Y., Colangelo, M., Collins, S., Collamore, A., Cooke, P.,
Dearellano, K., Dewar, K., Dodge, S., Domino, M., Doyle, M.,
Fenster, J., Ferreira, P., Fitzlugh, W., Forrest, C., Gage, D.,
Galsgott, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L.,
Grand-Pierre, N., Grant, G., Hagos, B., Heatard, A., Horton, L.,
Howard, J. C., Iliev, I., Johnson, R., Jones, C., Kann, L., Karatas, A.,
Klein, J., Landers, T., Laroque, K., Lechoczky, R., Levine, R.,
Lieu, C., Liu, G., Locke, K., MacDonald, P., Margus, N., McCarthy, M.,
Mcwan, P., McGurk, A., McKernan, K., McPheters, R., Meldrum, J.,
Meneus, L., Mihova, T., Miranda, C., Mlenga, V., Morrow, J., Naylor, J.,
Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliva, T. M.,
Peterson, K., Pierre, N., Pisanli, C., Pollara, V., Raymond, C.,
Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S.,
Severy, P., Spencer, B., Strange-Thomann, N., Stojanovic, N.,
Subramanian, A., Talamas, J., Teefaye, S., Theodore, J., Tirrell, A.,
Travers, M., Triggillo, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B.,
Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zimmer, A. and
Zody, M.

TITLE Direct Submission
JOURNAL Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT On Jun 7, 2000 this sequence version replaced gi:6970573.

Smit, A.F.A. & Green, P. (1996-1997)

----- Genome Center

Center code: WIBR

Contact: sequence_submissions@genome.wi.mit.edu

Center project name: L6363

----- Summary Statistics -----

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phipp, version 0.500/3.1
Consensus quality: 135238 bases at least

consensus quality: 141969 bases at least
consensus quality: 146244 bases at least

Insert size: 150000; agarose- μ p
Insert size: 151322; sum-of-contrasts

Quality coverage: 4.0 in Q20 bases; agarose-1p
Quality coverage: 3.9 in Q20 bases; sum-of-contrigs

NOTE: This is a 'working draft' sequence. It currently consists of 22 contrigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contrigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1152: contrig of 1152 bp in length
1153 1252: gap of 100 bp
1253 2252: contrig of 1000 bp in length
2253 2352: gap of 100 bp
2353 3503: contrig of 1151 bp in length
3504 3603: gap of 100 bp
3604 4658: contrig of 1055 bp in length
4659 4758: gap of 100 bp
4759 6059: contrig of 1301 bp in length
6060 6159: gap of 100 bp
6160 6832: contrig of 673 bp in length
6833 6932: gap of 100 bp
6933 9547: contrig of 2615 bp in length
9548 9647: gap of 100 bp
9648 11267: contrig of 1620 bp in length
11268 11367: gap of 100 bp
11368 14730: contrig of 3363 bp in length
14731 14830: gap of 100 bp
14831 18501: contrig of 3671 bp in length
18502 18601: gap of 100 bp
18602 23876: contrig of 5275 bp in length
23877 23976: gap of 100 bp
23977 28125: contrig of 4149 bp in length
28126 28225: gap of 100 bp
28226 33384: contrig of 5159 bp in length
33385 33484: gap of 100 bp
33485 37950: contrig of 4466 bp in length
37951 38050: gap of 100 bp
38051 43278: contrig of 5228 bp in length
43279 43378: gap of 100 bp
43379 49641: contrig of 6263 bp in length
49642 49741: gap of 100 bp
49742 57239: contrig of 7498 bp in length
57240 57339: gap of 100 bp
57340 65126: contrig of 7787 bp in length
65127 65226: gap of 100 bp
65227 74642: contrig of 9416 bp in length
74643 74742: gap of 100 bp
74743 86960: contrig of 12218 bp in length
86961 87060: gap of 100 bp
115867 115886: contrig of 28826 bp in length
115887 115987: gap of 100 bp
115987 153422: contrig of 37436 bp in length.

FEATURES
source

1. 153422
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/map="6"
/clone="RP11-612M16"
/clone_lib="RP11 Human Male BAC"
1. 1152
/note="assembly_fragment"
misc_feature
1253. 2252
/note="assembly_fragment"
misc_feature
2353. 3503
/note="assembly_fragment"
misc_feature
3604. 4658
/note="assembly_fragment"
misc_feature
4759. 6059
/note="assembly_fragment"
misc_feature
6160. 6832

/note="assembly_fragment"
clone_end:17
vector_side:right"
misc_feature
6933. 9547
/note="assembly_fragment"
9648. 11267
/note="assembly_fragment"
11368. 14730
/note="assembly_fragment"
14831. 18501
/note="assembly_fragment"
18602. 23876
/note="assembly_fragment"
23977. 28125
/note="assembly_fragment"
28226. 33384
/note="assembly_fragment"
33485. 37950
/note="assembly_fragment"
38051. 43278
/note="assembly_fragment"
clone_end:SP6
vector_side:right"
misc_feature
43379. 49641
/note="assembly_fragment"
49742. 57239
/note="assembly_fragment"
57340. 65126
/note="assembly_fragment"
65227. 74642
/note="assembly_fragment"
74743. 86960
/note="assembly_fragment"
87061. 115886
/note="assembly_fragment"
115987. 153422
/note="assembly_fragment"
BASE COUNT 47446 a 29368 c 28599 g 45687 t 2102 others
ORIGIN

Query Match 57.3%; Score 63.6; DB 2; Length 153422;
Best Local Similarity 73.6%; Pred. No. 8.7e-11;
Matches 81; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTGGCTTCAGAGATCTTTTATCTTTCAGAGACTTGCGGCG 60
DB 49270 ATGCGAGACCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 49329
QY 61 GAGTATGTAACGTCGCGGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 110
DB 49330 GAGTATGTAACGTCGCGGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 49379

RESULT 10
AC109592/c 86314 bp DNA linear PRI 13-MAR-2002
LOCUS Homo sapiens BAC clone RP11-750022 from 4, complete sequence.
DEFINITION AC109592
ACCESSION AC109592
VERSION AC109592.3 GI:19310357
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homini; Homo.
REFERENCE 1 (bases 1 to 86314)
Sulston, J.E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
PUBMED 9847074
REFERENCE 2 (bases 1 to 86314)
Buatsi, D. and Kozlowicz, A.
AUTHORS The sequence of Homo sapiens BAC clone RP11-750022
TITLE

JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (05-FEB-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
REFERENCE 4 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (06-MAR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
REFERENCE 5 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (09-MAR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
REFERENCE 6 (bases 1 to 86314)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (13-MAR-2002) Department of Genetics, Washington University 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Mar 9, 2002 this sequence version replaced gl:18642965.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: saplens@watson.wustl.edu
----- Summary Statistics
Center project name: H_NH0750022

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see http://genome.wustl.edu/gsc

SOURCE INFORMATION:
The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Moon,P.Y., Zhao,B., Frengen,E., Tateno,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (http://www.resgen.com) or Pieter de Jong and coworkers at http://www.chori.org
VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-510D4, 2000 bp overlap; the clone sequenced to the right is RP11-36611, 2000 bp overlap. Actual start of this clone is at base position 149897 of RP11-510D4; actual end is at base position 44359 of RP11-36611.

FEATURES
Source
1..86314
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="4"

/map="4"
/clone="RP11-750022"
/clone_lib="RPCI-11"
2..160
/rpt_family="Alu"
repeat_region
256..655
/rpt_family="L1"
repeat_region
954..1182
/rpt_family="L1"
repeat_region
1216..1472
/rpt_family="L1"
repeat_region
1563..1876
/rpt_family="L1"
repeat_region
2896..3255
/rpt_family="MaLR"
repeat_region
3256..4669
/rpt_family="MaLR"
repeat_region
4670..4969
/rpt_family="MaLR"
repeat_region
10088..10184
/rpt_family="MaLR"
repeat_region
11452..11617
/rpt_family="CRL"
repeat_region
12716..12930
/rpt_family="CRL"
repeat_region
13258..13395
/rpt_family="MIR"
repeat_region
14378..14415
/rpt_family="AT-rich"
repeat_region
14817..14883
/rpt_family="Alu"
repeat_region
15540..15842
/note="match to EST AI023549 (NID:g3238593) ov79f12.s1"
misc_feature
21639..21945
/rpt_family="L1"
repeat_region
23095..23142
/rpt_family="(TG)n"
repeat_region
24837..24987
/rpt_family="ERV1"
repeat_region
24994..26519
/rpt_family="ERV1"
repeat_region
26550..26833
/rpt_family="ERV1"
repeat_region
26834..26918
/rpt_family="(TA)n"
repeat_region
26938..26973
/rpt_family="(GA)n"
repeat_region
26979..27252
/rpt_family="Alu"
repeat_region
27546..27618
/rpt_family="MaLR"
repeat_region
27984..28617
/rpt_family="L1"
repeat_region
28624..28942
/rpt_family="Alu"
repeat_region
29073..29150
/rpt_family="MER53"
repeat_region
29081..29158
/rpt_family="MER53"
repeat_region
29199..29562
/rpt_family="L1"
repeat_region
29563..30039
/rpt_family="L1"
repeat_region
30313..31555
/rpt_family="L1"
repeat_region
31554..31738
/rpt_family="L1"
repeat_region
31739..31786
/rpt_family="AT-rich"
repeat_region
34024..34352
/rpt_family="Alu"
repeat_region
38180..38776
/rpt_family="L1"

```

repeat_region      3867..39139
                    /rpl_family="L1"
repeat_region      39172..39448
                    /rpl_family="L1"
repeat_region      39523..39821
                    /rpl_family="Alu"
repeat_region      40157..40231
                    /rpl_family="AT-rich"
repeat_region      40341..40597
                    /rpl_family="L1"
repeat_region      40604..40670
                    /rpl_family="(TG)n"
repeat_region      40758..40966
                    /rpl_family="L1"
repeat_region      41763..41802
                    /rpl_family="AT-rich"
repeat_region      42632..42724
                    /rpl_family="(CATATA)n"
repeat_region      42858..42895
                    /rpl_family="L1"
repeat_region      42901..43787
                    /rpl_family="L1"
repeat_region      43786..45120
                    /rpl_family="L1"
repeat_region      45123..45419
                    /rpl_family="MALR"
repeat_region      45531..47107
                    /rpl_family="MALR"
repeat_region      47108..47262
                    /rpl_family="MALR"
repeat_region      47263..48876
                    /rpl_family="L1"
repeat_region      51158..51280
                    /rpl_family="Achobo"
repeat_region      51281..51586
                    /rpl_family="L1"

```

Query Match 57.1%; Score 63.4; DB 9; Length 86314;
 Best Local Similarity 82.0%; Pred. No. 1e-10;
 Matches 73; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

```

OY 23 AGCAATCTTTTCATCTTTCAGAGACCTTCGGCGGAGAGATGATAACTCTCGGCTC 82
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 32282 AGGATGAGATCTTCTCCCTTCGGCGGAGATCTTCGGCTGAGATGATAAATCTTGGGCTC 32223
OY 83 TCTGTGTGTGCTCGAGTGGCTCTCTACT 111
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 32222 TCTGTGTGTGCCAGTGGCTCTCTGCT 32194

```

```

RESULT 11
AC021378/c 159475 bp DNA linear HTG 26-MAY-2000
LOCUS Homo sapiens clone RP11-26G6, WORKING DRAFT SEQUENCE, 10 unordered
DEFINITION AC021378
ACCESSION AC021378
VERSION AC021378.4 GI:8072570
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 159475)
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.
JOURNAL Homo sapiens, clone RP11-26G6
TITLE Unpublished
REFERENCE 2 (bases 1 to 159475)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
JOURNAL Anderson,S., Baldwin,J., Barna,N., Beckert,R., Beda,F.,
Boguslavsky,L., Bouckhailler,B., Brown,A., Burkett,G., Castle,A.,
Chepel,I., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
DeRellano,K., Dewar,K., Domino,M., Doyle,M., Fenestor,J.,
Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,

```

TTITLE JOURNAL COMMENT

Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
 Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatsis,A., Klein,J.,
 Landers,T., Lehoczeky,J., Levine,R., Lien,C., Liu,G., Locke,K.,
 MacDonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
 McPheeters,R., Meldrum,J., Menius,L., Morrow,J., Naylor,J.,
 Norman,C.H., O'Connor,T., O'Donnell,P., Oliver,T.M., Peterson,K.,
 Pierre,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
 Roy,A., Santos,R., Severy,P., Spencer,B., Strange-Thomann,N.,
 Stojanovic,N., Suramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
 Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
 Zimmer,A., and Zody,M.
 Direct Submission
 Submitted (16-JAN-2000) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On May 25, 2000 this sequence version, replaced gi:7637277.
 All repeats were identified using RepeatMasker:
 Smit,A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: MIBR
 Web site: http://www.seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 Project Information
 Center project name: L4633
 Center clone name: 26.G.6

----- Summary Statistics -----
 Sequencing vector: M13; M77815; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 150152 bases at least Q40
 Consensus quality: 15379 bases at least Q30
 Consensus quality: 157415 bases at least Q20
 Insert size: 157000; agarose-fp
 Insert size: 158575; sum-of-contigs
 Quality coverage: 4.2 in Q20 bases; agarose-fp
 Quality coverage: 4.2 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 10 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

```

1 2315: contig of 2315 bp in length
2316 2415: gap of 100 bp
2416 9041: contig of 6626 bp in length
9042 9141: gap of 100 bp
9142 17359: contig of 8218 bp in length
17360 17459: gap of 100 bp
17460 26067: contig of 8608 bp in length
26068 26167: gap of 100 bp
26168 40876: contig of 14709 bp in length
40877 40976: gap of 100 bp
40977 56845: contig of 15869 bp in length
56846 56945: gap of 100 bp
56946 74524: contig of 17579 bp in length
74525 74624: gap of 100 bp
74625 94662: contig of 20038 bp in length
94663 94762: gap of 100 bp
94763 126925: contig of 32163 bp in length
126926 127025: gap of 100 bp
127026 159475: contig of 32450 bp in length.

```

FEATURES

source
 1..159475 "Homo sapiens"
 /db_xref="taxon:9606"
 /clone="RP11-26G6"
 /clone_lib="RPC1-11 Human Male BAC"
 1..2315
 /note="assembly_fragment
 misc_feature

```
misc_feature      clone_end:SP6
                  vector_side:left"
                  2416..9041
                  /note="assembly_fragment"
misc_feature      9142..17359
                  /note="assembly_fragment"
misc_feature      17460..26067
                  /note="assembly_fragment"
misc_feature      26168..40876
                  /note="assembly_fragment"
misc_feature      40977..56845
                  /note="assembly_fragment"
misc_feature      56946..74524
                  /note="assembly_fragment"
misc_feature      74625..94662
                  /note="assembly_fragment"
misc_feature      94763..126925
                  /note="assembly_fragment"
misc_feature      127026..159475
                  /note="assembly_fragment"
misc_feature      /note="assembly_fragment"
                  clone_end:T7
                  vector_side:left"
BASE COUNT      52037 a 27831 c 28355 g 50348 t 904 others
ORIGIN
```

```
Query Match      57.1% Score 63.4: DB 2: Length 159475;
Best Local Similarity 82.0% Pred. NO. 1e-10; Index 0; Gaps 0;
Matches 73: Conservative 0; Mismatches 16; Gaps 0;

QY 23 AGAATCTTTTTCATCTTTCGACGACTTCTGGGCGCGAGTATGTAACCTCTGGGTC 82
      |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 45765 AGAATGATCTTTCGCTGCGGGAATCTGGGCGCGAGTATGTAACCTCTGGGTC 45706
      |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
QY 83 TCTGTGTGCTGCTGAGTGGCTGCTCTACT 111
      |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 45705 TCTGTGTGCTGCTGAGTGGCTGCTCTACT 45677
```

```
RESULT 12
AL157937/c      153783 bp DNA linear PRI 04-APR-2001
LOCUS          Human DNA sequence from clone RP11-408N14 on chromosome 9 Contains
DEFINITION     STS and GSSs, complete sequence.
ACCESSION     AL157937
VERSION       AL157937.21 GI:11121006
KEYWORDS      HTG.
SOURCE        human.
ORGANISM      Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE     1 (bases 1 to 153783)
AUTHORS       Kimberley A.
TITLE         Direct Submission
JOURNAL       Submitted (26-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
              CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
              requests: clonerequests@sanger.ac.uk
              On Nov 8, 2000 this sequence version replaced gi:11071307.
              During sequence assembly data is compared from overlapping clones.
              Where differences are found these are annotated as variations
              together with a note of the overlapping clone name. Note that the
              validation annotation may not be found in the sequence submission
              corresponding to the overlapping clone, as we submit sequences with
              only a small overlap as described above.
              The following abbreviations are used to associate primary accessions
              numbers given in the feature table with their source databases:
              Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information
              http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
              was generated from part of bacterial clone contigs of human
              chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping
              Group. Further information can be found at
              http://www.sanger.ac.uk/HGP/Chr9
              IMPORTANT: This sequence is not the entire insert of clone
```

RP11-408N14 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone RP11-408N14 is at 1 in this sequence. The true left end of clone RP11-44115 is at 13364 in this sequence. The true right end of clone RP11-145E5 is at 29400 in this sequence. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. RP11-408N14 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

FEATURES

source

```
repeat_region    /note="MIR repeat: matches 21..190 of consensus"
repeat_region    /note="L1M4 repeat: matches 4921..5372 of consensus"
repeat_region    /note="L1M4 repeat: matches 1..304 of consensus"
repeat_region    /note="L1M5 repeat: matches 7249..7897 of consensus"
repeat_region    /note="L1M5 repeat: matches 2401..2642 of consensus"
repeat_region    /note="L1M5 repeat: matches 6983..7227 of consensus"
repeat_region    /note="23 copies 2 mer tg 93% conserved"
repeat_region    /note="MIR repeat: matches 5..241 of consensus"
repeat_region    /note="MIR repeat: matches 5093..5394 of consensus"
repeat_region    /note="match: GSS: Em:A0823856"
repeat_region    /note="L2 repeat: matches 2164..2262 of consensus"
repeat_region    /note="L2 repeat: matches 5537..5881 of consensus"
repeat_region    /note="MLT2FA repeat: matches 5..324 of consensus"
repeat_region    /note="match: GSS: Em:A0137344"
repeat_region    /note="match: GSS: Em:B91342"
repeat_region    /note="match: GSS: Em:A0217550"
repeat_region    /note="L2 repeat: matches 1..297 of consensus"
repeat_region    /note="L2 repeat: matches 2651..2694 of consensus"
repeat_region    /note="L2 repeat: matches 1..309 of consensus"
repeat_region    /note="L2 repeat: matches 2576..2651 of consensus"
repeat_region    /note="L2 repeat: matches 20..204 of consensus"
repeat_region    /note="MIR repeat: matches 29..213 of consensus"
repeat_region    /note="match: GSS: Em:A0771822"
repeat_region    /note="36 copies 2 mer tg 79% conserved"
repeat_region    /note="MIR repeat: matches 139..262 of consensus"
repeat_region    /note="L1P45 repeat: matches 76..6143 of consensus"
repeat_region    /note="L1M4 repeat: matches 7641..7835 of consensus"
```


repeat_region	18520..20148	/note="L1P12 repeat: matches 1421..1009 of consensus"
repeat_region	20127..21370	/note="L1P11 repeat: matches 921..2161 of consensus"
repeat_region	21372..21581	/note="L1P11 repeat: matches 2262..6165 of consensus"
repeat_region	25293..25713	/note="L1P10 repeat: matches 5655..6083 of consensus"
repeat_region	26120..26412	/note="L1P12 repeat: matches 1541..1650 of consensus"
misc_feature	26198..26577	/note="match: GSS: Em:AQ147782"
repeat_region	26660..26962	/note="L1P12 repeat: matches 1..300 of consensus"
misc_feature	complement(27055..27448)	/note="match: STS: Em:HSJ37G11"
repeat_region	27407..31040	/note="L1P12 repeat: matches 2505..6164 of consensus"
repeat_region	31031..31138	/note="L1P repeat: matches 1541..1650 of consensus"
repeat_region	31144..31497	/note="HEB repeat: matches 1..364 of consensus"
repeat_region	31498..33130	/note="HEB-INTERNAL repeat: matches 1..1580 of consensus"
repeat_region	33131..33479	/note="HEB repeat: matches 1..359 of consensus"
repeat_region	33485..35130	/note="L1P13 repeat: matches 248..1557 of consensus"
repeat_region	36624..36669	/note="23 copies 2 mer ta 76% conserved"
repeat_region	36680..36897	/note="LMC repeat: matches 6404..6623 of consensus"
repeat_region	37335..37508	/note="37 copies 2 mer ag 75% conserved"
repeat_region	38923..39151	/note="Charlie4 repeat: matches 231..1926 of consensus"
misc_feature	complement(39281..39766)	/note="match: GSS: Em:AQ894025"
repeat_region	40317..40540	/note="MR repeat: matches 20..234 of consensus"
repeat_region	40603..40726	/note="12 repeat: matches 2587..2710 of consensus"
repeat_region	43871..43952	/note="12 repeat: matches 2673..2750 of consensus"
repeat_region	43962..44259	/note="Alu repeat: matches 1..299 of consensus"
repeat_region	45076..45280	/note="MR repeat: matches 2..205 of consensus"
repeat_region	45413..45588	/note="MR repeat: matches 1..205 of consensus"
misc_feature	complement(46149..46586)	/note="match: GSS: Em:AQ343245"
repeat_region	47817..47938	/note="12 repeat: matches 2627..2749 of consensus"
repeat_region	48349..48579	/note="L1P11 repeat: matches 1..220 of consensus"
repeat_region	50262..50664	/note="12 repeat: matches 2339..2749 of consensus"
repeat_region	50674..52080	/note="L1M1 repeat: matches 1389..30 of consensus"
repeat_region	52084..52853	/note="L1P12 repeat: matches 1541..1650 of consensus"
repeat_region	54498..60658	/note="L1P12 repeat: matches 1..6146 of consensus"
repeat_region	61367..61560	/note="L1P12 repeat: matches 346..541 of consensus"
repeat_region	62305..62430	/note="63 copies 2 mer tt 59% conserved"
repeat_region	63089..63699	/note="L1P12 repeat: matches 10..540 of consensus"
repeat_region	63630..63887	/note="Alu repeat: matches 7..263 of consensus"

```

repeat_region 63888..64059 /note="A1u5g repeat: matches 134..303 of consensus"
repeat_region 64060..64352 /note="A1u5g repeat: matches 1..295 of consensus"
repeat_region 64353..64485 /note="A1u5g repeat: matches 1..134 of consensus"
repeat_region 64486..64834 /note="A1u5g repeat: matches 1..291 of consensus"
repeat_region 64816..64938 /note="L1P1A12 repeat: matches 1761..1897 of consensus"
repeat_region 64939..65350 /note="L1P1A10 repeat: matches 1..608 of consensus"
repeat_region 65351..67266 /note="L1R14B repeat: matches 1894..3818 of consensus"
repeat_region 67317..69162 /note="L1P1A10 repeat: matches 3793..5643 of consensus"
repeat_region 69122..69167 /note="L1P1A10 repeat: matches 5616..5662 of consensus"
repeat_region 69216..69754 /note="L1P1A repeat: matches 5610..6165 of consensus"
repeat_region 70283..70369 /note="L1P1A10 repeat: matches 5610..6165 of consensus"

```

Query Match	56.8%;	Score 63;	DB 9;	Length 153783;
Best Local Similarity	73.0%;	Pred. No. 1.4e-10;		
Matches	81;	Conservative	0;	Mismatches 30; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTTTCCTTGACAGATTCTTTTCATCTTTTGACAGGAATCTTGGGCCG 60
||| ||| |,||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 18987 AATGACAGACAATCCTGCTCTGTCTGAGTTGCAGACACCCTTGTGCGGGCAATCTGGGCCG 18928

Qy 61 GAGTATGTAAACTCCTCGGCTCTCTGTGTGTGCTGAGTGGCTCTACT 111
|||||
Db 18927 GAGTATGTAAAGCTCCTGGTCTCTGTGCATACCTGAGCAGCTGCTCTGCT 18877
|||||

RESULT 13	
AC021025/c	
ID AC021025	standard; DNA; HTG; 123779 BP.

XX	
AC	AC021025;
XX	
SV	AC021025.9
XX	
DT	14-JAN-2000 (Rel. 62, Created)
	08-AUG-2000 (Rel. 64, Last updated, Version 11)

AA Homo sapiens chromosome 3 clone RP11-79K17, WORKING DRAFT SEQUENCE, 10
 DE
 DE unordered pieces.
 XX
 TW HMG; HTGS_DRAFT; HTGS_PHASE1.

05 Homo sapiens (human)
0C Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia
0C Eutheria; primates; Catarrhini; Hominidae; Homo.

RA Muzny D.M., Adams C., Bailey M., Barbara J., Blankenburg K., Bodota B.,
 RP 1-123779
 KIN 143

RA Ding Y., Domani-Rasniun N., Dugan-Rocha S., Durbin K.V., Fernandez C., Ferraguto D., Forcum-Tansey J., Frantz P., Ganesh R., Gorrell J.H., RA Gorrell L.L., Guevara W., Harris K., Hernandez J., Hodgson A., Hogues M.,

KA Liu J., Liu W., Logan C., Lozaco R.J., Lu J., Lucier R., Martin R.,
RA Martinez C., McLeod M.P., Mei G., Morgan M., Morris S., Nash S., Nelson A.,
RA Nguyen R., Nguyen N., Nguyen S., Oswald G., Parish B., Paxton S., Payton B.

KA Tabor H., Taylor I., Vasquez L., Vinson R., VO V., Wampan M.,
RA Watlington S., Weinstein G., Weinstein I.R., Williamson A., Worley K.,
RA Wren J., Wrensfeld G., Yu W., Zhou X., Nelson D., Gibbs R.;

RT "Direct Submission";
RL Unpublished.
XX [2]
RN 1-123779
RA Morley K.C.;
RT Submitted (12-JAN-2000) to the EMBL/GenBank/DBJ databases.
RL Human Genome Sequencing Center, Department of Molecular and Human Genetics,
Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
XX
CC On Aug 7, 2000 this sequence version replaced g1:8700003.
CC ----- Genome Center
CC Center: Baylor College of Medicine
CC Center code: BCM
CC Web site: <http://www.hgsc.bcm.tmc.edu/>
CC Contact: hgsc-help@bcm.tmc.edu
CC ----- Project Information
CC Center project name: HMXB
CC Center clone name: RP11-79K17
CC ----- Summary Statistics
CC Sequencing vector: M13; L08821
CC Chemistry: Dye-terminator Big Dye; 100% of reads
CC Assembly program: Phrap; version 0.990329
CC Consensus quality: 112332 bases at least Q40
CC Consensus quality: 117272 bases at least Q30
CC Consensus quality: 119789 bases at least Q20
CC Estimated insert size: 121963; sum-of-coverage estimation
CC Quality coverage: 0x in Q20 bases; agarose-gel estimation
CC Quality coverage: 5.1x in Q20 bases; sum-of-coverage estimation
CC -----
CC * NOTE: Estimated insert size may differ from sequence length
CC * (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
CC * NOTE: This is a 'working draft' sequence. It currently
CC * consists of 10 contigs. The true order of the pieces
CC * is not known and their order in this sequence record is
CC * arbitrary. Gaps between the contigs are represented as
CC * runs of N, but the exact sizes of the gaps are unknown.
CC * This record will be updated with the finished sequence
CC * as soon as it is available and the accession number will
CC * be preserved.
CC 1
CC 35931: contig of 35931 bp in length
CC 35932 36031: gap of unknown length
CC 36032 60754: contig of 24723 bp in length
CC 60755 60854: gap of unknown length
CC 60855 71667: contig of 10813 bp in length
CC 71668 71768: gap of unknown length
CC 71769 84826: contig of 13059 bp in length
CC 84827 84926: gap of unknown length
CC 84927 96721: contig of 11795 bp in length
CC 96722 96821: gap of unknown length
CC 96822 105594: contig of 8773 bp in length
CC 105595 105694: gap of unknown length
CC 105695 112907: contig of 7213 bp in length
CC 112908 113007: gap of unknown length
CC 113008 120198: contig of 7191 bp in length
CC 120199 120298: gap of unknown length
CC 120299 122220: contig of 1922 bp in length
CC 122221 122321: gap of unknown length
CC 122322 123779: contig of 1459 bp in length.
XX
XX
FH Key Location/Qualifiers
FH
FT source 1. 123779
FT /chromosome="3"
FT /db_xref="taxon:9606"
FT /organism="Homo sapiens"
FT /clone="RP11-79K17"
XX
SQ Sequence 123779 BP; 37802 A; 24017 C; 23437 G; 37612 T; 911 other;
Query Match 56.2%; Score 62.4; DB 30; Length 123779;
Best Local Similarity 81.8%; Pred. NO. 2.2e-10;

Matches 72; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
QY 24 GCATCTTTTTCATCTTTCGACAGGACTTCGGGCGCGAGATATGTAACCTCCGGGCT 83
DB 3128 GGATGATCTCCACACCTGCTGAGATCTGAGGAGTATGATCAAACTCTGGGCT 3069
QY 84 CTGTGTGCTGAGTGGCTGCTACT 111
DB 3068 CTGTGTGCTGAGTGGCTGCTACT 3041
RESULT 14
AC076969 128118 bp DNA linear HTG 15-OCT-2001
LOCUS AC076969
DEFINITION Homo sapiens chromosome 3 clone RP11-79K12, WORKING DRAFT SEQUENCE,
14 unordered pieces.
ACCESSION AC076969
VERSION AC076969.6 GI:16117967
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS
1 (bases 1 to 128118)
Muzny,D.M., Adams,C., Adio-Oduola,B., Alt-osman,F.R., Allen,C.,
Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Banks,T., Barbata,J.,
Benton,J., Blamege,K., Blankenburg,K., Bonnin,D., Bouck,J.,
Bowie,S., Brileva,M., Brown,M., Bryant,N.P., Buhay,C.,
Burke,P., Burkett,C., Burrell,K.L., Byrd,N.C., Cartron,T.F.,
Carrier,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,
Chen,Z., Chowdhury,I., Christopoulos,C., Cleveland,C.D., Cox,C.,
Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,
Davy,Carrill,L., Dederich,D.A., Delaney,K.R., Delgado,O.,
Denn,A.L., Ding,Y., Dinh,H.H., Doutharte,K.J., Draper,H.,
Dugan-Rocha,S., Durbin,K.J., Earhart,C., Edgar,D., Edwards,C.C.,
Ellis,J.C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,
Foster,P., Frantz,P., Gabisi,A., Guevara,W., Gunaratne,P., Hale,S.,
Hamilton,K., Harris,C., Harris,K., Hart,M., Hayla,P., Hawes,A.,
Hernandez,J., Hernandez,O., Hodgson,A., Hoques,M., Holloway,C.,
Hollins,B., Homsf., Howard,S., Huber,J., Hulky,S., Hume,J.,
Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S.,
Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J.,
Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,
Lewis,L., Li,J., Li,Z., Lichtenarge,O., Liu,C., Liu,J., Liu,W.,
Louisleged,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,
Ma,J., Maheshwari,M., Mapa,P., Martin,R., Martindale,A.,
Martinez,E., Massey,E., Mawhiney,E., McLeod,M.P., Meador,M.,
Mei,G., Metzker,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K.,
Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,N.,
Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokenko,S.,
Ogunu,M., Okunolu,G., Oragunye,N., Oviedo,R., Pace,A., Payton,B.,
Peery,J., Perez,L., Peters,L., Pickens,R., Pritus,E., Pu,L.L.,
Quiles,M., Ren,Y., Rivers,M., Rojas,A., Rojuben,I., Rolfe,M.,
Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shooshitari,N.,
Stison,I., Sodergren,E., Sonalke,T., Sparks,A., Stanley,H.,
Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K.,
Tang,H., Tansey,J., Taylor,C., Taylor,T., Talford,B., Thomas,N.,
Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalob,D., Vinson,R.,
Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
Washington,S., Williams,G., Williamson,A., Wlezyk,R., Wooden,S.,
Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 128118)
MORLEY K.C.
Direct Submission
Submitted (01-AUG-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Oct 14, 2001 this sequence version replaced g1:10047573.
----- Genome Center

covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bp) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

NsII		BglII		EcoRI	
SeqDerMap	FngPrnt	SeqDerMap	FngPrnt	SeqDerMap	FngPrnt
-----	-----	-----	-----	-----	-----
12049	12136	4077	4013	8696	8902
-----	-----	-----	-----	-----	-----
579	<800	2067	2145	6	<800
-----	-----	-----	-----	-----	-----
1056	1065	9698	9932	1323	1337
-----	-----	-----	-----	-----	-----
13085	13126	3729	3734	1024	961
-----	-----	-----	-----	-----	-----
10944	10836	1632	1616	8421	8408
-----	-----	-----	-----	-----	-----
581	<800	8069	8139	1114	1126
-----	-----	-----	-----	-----	-----
882	887	1507	1485	3538	3534
-----	-----	-----	-----	-----	-----
8435	8467	11939	11871	6051	6022
-----	-----	-----	-----	-----	-----
3924	4103	133	<800	5129	5048
-----	-----	-----	-----	-----	-----
9842	9759	2135	2145	58	<800
-----	-----	-----	-----	-----	-----
4401	4377	78	<800	1895	1903
-----	-----	-----	-----	-----	-----
4458	4377	5873	5861	799	<800
-----	-----	-----	-----	-----	-----
4587	4522	896	902	858	885
-----	-----	-----	-----	-----	-----
6048	5947	3312	3356	931	885
-----	-----	-----	-----	-----	-----
1395	1385	1505	1485	859	885
-----	-----	-----	-----	-----	-----
3059	3071	6694	6716	3934	3936
-----	-----	-----	-----	-----	-----
3911	3861	10032	9932	1065	1058
-----	-----	-----	-----	-----	-----
138	<800	2168	2145	2629	2723
-----	-----	-----	-----	-----	-----
50	<800	1794	1785	859	885
-----	-----	-----	-----	-----	-----
582	<800	227	<800	6855	6888
-----	-----	-----	-----	-----	-----
2225	2229	4002	4013	7821	7819
-----	-----	-----	-----	-----	-----
984	967	2714	2726	7210	7201
-----	-----	-----	-----	-----	-----
887	887	5628	5624	6992	6888
-----	-----	-----	-----	-----	-----
1042	1065	743	758	1204	1224
-----	-----	-----	-----	-----	-----
2783	2808	3045	2963	2722	2723
-----	-----	-----	-----	-----	-----
7074	7158	1288	1262	321	<800
-----	-----	-----	-----	-----	-----
4087	3861	967	973	1066	1058

7976	7926	461	<800	470	<800
-----	-----	-----	-----	-----	-----
1233	1199	4649	4618	3055	3072
-----	-----	-----	-----	-----	-----
3875	3861	5289	5417	10961	10848
-----	-----	-----	-----	-----	-----
27	<800	1098	1095	6725	6888
-----	-----	-----	-----	-----	-----
1137	1065	798	809	1896	1903
-----	-----	-----	-----	-----	-----
750	749	111	<800	8682	8666
-----	-----	-----	-----	-----	-----
214	<800	7644	7609	4438	4342
-----	-----	-----	-----	-----	-----
836	887	10084	9932	514	<800
-----	-----	-----	-----	-----	-----
1670	1651	3641	3589	3077	3072
-----	-----	-----	-----	-----	-----
2650	2609	7555	7609	1371	1337
-----	-----	-----	-----	-----	-----
1890	1872	-----	-----	4185	4087
-----	-----	-----	-----	-----	-----
4615	4522	-----	-----	719	<800
-----	-----	-----	-----	-----	-----
1321	1273	-----	-----	4130	4087
-----	-----	-----	-----	-----	-----
-----	-----	-----	-----	367	<800
-----	-----	-----	-----	2979	3072
-----	-----	-----	-----	-----	-----
-----	-----	-----	-----	333	<800

FEATURES

source 1. 128583
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="3"
 /clone="RP11-79K17"
 /clone_lib="RPC1 human BAC library 11"
 BASE COUNT 37777 a 24752 c 24789 g 41265 t
 ORIGIN

Query Match 56.2%; Score 62.4; DB 9; Length 128583;

Best Local Similarity 81.8%; Pred. NO. 2.2e-10; Mismatches 16; Indels 0; Gaps 0;

Matches 72; Conservative 0;

QY 24 GGATTCCTTTTCATCTTGGCAGGACCTCTGGGGCCGGAGTATGTAACCTCGGTCT 83

DB 27328 GGATGATCTCCACCTTCTGAGAACTCTGGGGTGGAGTATGCAAACTCTGGTCT 27269

QY 84 CTGTGTGCTGCTGAGTGGCTGCTACT 111

DB 27268 CTGTGTGCTGCTGAGTGGCTGCTACT 27241

Search completed: April 25, 2003, 02:17:54
 Job time : 1357 secs

GenCore version 5.1.4-p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:52:59 ; Search time 150 seconds
(without alignments)
1666.479 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atggttgatccttgcctt.....gcctgagtgcctctact 111

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 2185239 segs, 112599159 residues

1 number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_101002.*

- 1: /SID52/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
- 2: /SID52/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
- 3: /SID52/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
- 4: /SID52/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
- 5: /SID52/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
- 6: /SID52/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
- 7: /SID52/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
- 8: /SID52/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
- 9: /SID52/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
- 10: /SID52/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
- 11: /SID52/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
- 12: /SID52/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
- 13: /SID52/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
- 14: /SID52/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
- 15: /SID52/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
- 16: /SID52/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
- 17: /SID52/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
- 18: /SID52/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
- 19: /SID52/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
- 20: /SID52/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
- 21: /SID52/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
- 22: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
- 23: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
- 24: /SID52/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	447	21	AAC03794 Human secreted pro
2	111	100.0	447	21	AAZ42680 Human 5' EST isola
3	58.2	52.4	349	22	AAE5886 Novel human polynu
4	58.2	52.4	570	22	ABA63453 Human foetal liver
5	58.2	52.4	570	22	ABA30652 Probe #9118 for ge
6	58.2	52.4	570	22	AAK11985 Human brain expres
7	58.2	52.4	570	22	AAK37688 Human bone marrow
8	58.2	52.4	570	22	AAI18447 Probe #8380 for ge
9	58.2	52.4	570	22	AAI43563 Probe #12249 used

C	10	58.2	52.4	570	24	ABS11680 Human genome-deriv
C	11	56.4	50.8	128600	24	ABK83461 Human cDNA differe
C	12	55	49.5	1982	21	AAC68089 Human secreted pro
C	13	52.8	47.6	609	22	AAH98980 Human EST-derived
C	14	50.8	45.8	1299	22	AAH91827 DNA encoding novel
C	15	46	41.4	807	23	AAH64647 DNA encoding novel
C	16	46	41.4	843	23	AAH79254 DNA encoding novel
C	17	44	39.6	1996	22	ABA18896 Human nervous syst
C	18	43.8	39.5	1909	23	AAH88434 DNA encoding novel
C	19	40.2	36.2	240	24	AAH25478 Human ORX polynuc
C	20	39	35.1	660	23	AAH73441 DNA encoding novel
C	21	38.8	35.0	162	22	ABA75762 Human foetal liver
C	22	38.8	35.0	519	22	ABA63320 DNA encoding novel
C	23	36.2	32.6	2787	23	AAH68019 DNA encoding novel
C	24	36.2	32.6	2787	23	AAH69828 DNA encoding novel
C	25	35.2	31.7	1173	23	AAH69200 DNA encoding novel
C	26	35.2	31.7	1173	23	AAH82303 DNA encoding novel
C	27	35.2	31.7	1173	23	AAH86064 DNA encoding novel
C	28	35.2	31.7	1291	23	AAH77610 DNA encoding novel
C	29	35.2	31.7	2097	23	AAH77443 DNA encoding novel
C	30	35.2	31.7	2569	23	AAH77685 DNA encoding novel
C	31	35.2	31.7	11534	24	ABL32342 Human immune syste
C	32	34.2	30.8	1278	23	AAH72474 DNA encoding novel
C	33	33.4	30.1	396	22	AAH74465 Human immune/haema
C	34	32.4	29.2	447	22	AAH16101 Human breast cance
C	35	32.4	29.2	447	22	AAH124945 Human breast cance
C	36	31.8	28.6	349980	22	AAH6431 Pyrococcus abyssi
C	37	31.8	28.6	349980	22	AAH41223 Human cDNA clone (
C	38	31	27.9	826	22	AAH04786 Interferon-pseudo-
C	39	30.8	27.7	3659	7	AAH60204 Human immune/haema
C	40	30.6	27.6	352	22	AAH59083 Human benign prost
C	41	29.2	26.3	369	24	ABK64361 Lung cancer relate
C	42	29.2	26.3	369	24	ABL61736 Lung cancer relate
C	43	29.2	26.3	369	24	ABL65612 Lung cancer relate
C	44	29.2	26.3	369	24	ABL66228 Human DNA for a no
C	45	29.2	26.3	1314	22	AAH31530

ALIGNMENTS

RESULT 1	AAC03794 standard; cDNA; 447 BP.
ID	XX AAC03794;
AC	XX AAC03794;
DT	XX 06-OCT-2000 (first entry)
XX	XX Human secreted protein 5' EST, SEQ ID NO: 3792.
DE	XX Human: 5' EST: expressed sequence tag; secreted protein; cDNA isolation;
XX	XX gene therapy; chromosome mapping; ss.
KW	XX Homo sapiens.
OS	XX EP1033401-A2.
XX	XX
PN	XX 06-SEP-2000.
PD	XX 21-FEB-2000; 2000EP-0200610.
XX	XX 26-FEB-1999; 99US-0122487.
PR	XX (GEST) GENSET.
PA	XX Dumas Milne Edwards J, Duclert A, Giordano J;
XX	XX WPI: 2000-500381/45.
PI	XX P-PDB; AAC03788.
XX	XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
DR	XX obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
PT	PT

PT diagnostic, forensic, gene therapy and chromosome mapping procedures -
XX
XX Claim 1: SEQ ID 3792; 71pp + CD-ROM; English.
XX
CC The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. An ORF has been identified within the
CC sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs
CC derived from 30 different tissues. EST sequences usually correspond
CC mainly to the 3' untranslated region (UTR) of the mRNA because they are
CC often obtained from oligo-dT primed cDNA libraries. Such ESTs are not
CC well suited for isolating cDNA sequences derived from the 5' ends of
CC mRNAs and even in those cases where longer cDNA sequences have been
CC obtained, the full 5' UTR is rarely included. 5' ESTs are derived from
CC mRNAs with intact 5' ends and can therefore be used to obtain full length
CC cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic,
CC gene therapy and chromosome mapping procedures. They are used to obtain
CC upstream regulatory sequences and to design expression and secretion
CC vectors.
XX
XX Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;
XX
XX Query Match 100.0%; Score 111; DB 21; Length 447;
XX Best Local Similarity 100.0%; Pred. No. 1.4e-27;
XX Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ATGGGTGATCTTTGCGCTGCGAGATCTTTTCATCTTTGACAGGACTTCTGGGCGC 60
DB 51 ATGGGTGATCTTTGCGCTGCGAGATCTTTTCATCTTTGACAGGACTTCTGGGCGC 110
OY 61 GAGTATGTAAACTCTGCGTCTGCTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTACT 111
DB 111 GAGTATGTAAACTCTGCGTCTGCTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTACT 161
RESULT 2
AAZ42680
ID AAZ42680 standard; cDNA; 447 BP.
XX
XX AAZ42680;
XX
DT 01-FEB-2000 (first entry)
XX
XX Human 5' EST isolated from a cDNA library SEQ ID NO:439.
XX
XX Human; 5' EST; expressed sequence tag; secreted protein; diagnosis;
XX gene therapy; chromosome mapping; upstream regulatory sequence;
XX forensic; location; development; protein synthesis; stability;
XX regulation; identification; ss.
XX
XX Homo sapiens.
XX
XX MO9953051-A2.
XX
XX 21-OCT-1999.
XX
XX 09-APR-1999; 99WO-IB00712.
XX
XX 09-APR-1998; 98US-0057719.
XX 28-APR-1998; 98US-0069047.
XX
XX (GIST) GENSET.
XX
XX Dunas Milne Edwards J, Duclert A, Giordano J;
XX
XX MPI: 2000-038446/03.
XX P-PSDB: AAY65066.
XX
XX Novel secreted protein 5' expressed sequence tag sequences used in
XX diagnostic, forensic, gene therapy, and chromosome mapping procedures
XX
XX Claim 1; Page 402; 837pp; English.
XX
XX AAZ42265 to AAZ43075 represent novel 5' expressed sequence tag (EST)

CC sequences, corresponding to human secreted proteins. AAY64651 to
CC AAY65438 represent the EST-related proteins corresponding to AAZ42265 to
CC AAZ43052. The 5' ESTs can be used for producing secreted human gene
CC products. They can be used to identify and isolate 5' untranslated
CC regions (UTRs) and upstream regulatory regions which control the
CC location, development stage, rate, and quantity of protein synthesis, as
CC well as stability of mRNA. The ESTs are also useful as probes for
CC chromosome mapping, and to obtain full length cDNA clones. The ESTs can
CC also be used in forensic procedures to identify individuals, or in
CC diagnostic procedures to identify individuals having genetic diseases
CC resulting from abnormal gene expression. The products may also be used in
CC gene therapy protocols. The nucleic acids encoding signal peptides can be
CC used for directing extracellular secretion of a polypeptide or the
CC insertion of a polypeptide into a membrane, or importing a polypeptide
CC into a cell. The proteins encoded by the EST sequences may be useful in
CC treating a variety of human conditions. Secreted proteins have
CC therapeutic value, and the identification of new secreted proteins is
CC valuable. AAZ42249 to AAZ4264 and AAY6464 to AAY64650 represent
CC sequences used in the exemplification of the present invention.
XX
XX
XX Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;
XX
XX Query Match 100.0%; Score 111; DB 21; Length 447;
XX Best Local Similarity 100.0%; Pred. No. 1.4e-27;
XX Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ATGGGTGATCTTTGCGCTGCGAGATCTTTTCATCTTTGACAGGACTTCTGGGCGC 60
DB 51 ATGGGTGATCTTTGCGCTGCGAGATCTTTTCATCTTTGACAGGACTTCTGGGCGC 110
OY 61 GAGTATGTAAACTCTGCGTCTGCTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTACT 111
DB 111 GAGTATGTAAACTCTGCGTCTGCTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTACT 161
RESULT 3
AAF65888/c
ID AAF65888 standard; cDNA; 349 BP.
XX
XX AAF65888;
XX
XX 09-APR-2001 (first entry)
XX
XX Novel human polynucleotide, SEQ ID NO: 1644.
XX
XX Human: cytostatic; gene therapy; colon cancer; prostate cancer;
XX breast cancer; lung cancer; cancer detection; ss.
XX
XX Homo sapiens.
XX
XX MO200102568-A2.
XX
XX 11-JAN-2001.
XX
XX 30-JUN-2000; 2000WO-US18374.
XX
XX 02-JUL-1999; 99US-0142310.
XX 02-JUL-1999; 99US-0142311.
XX
XX (CHIR) CHIRON CORP.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Williams LR, Escobedo J, Innis MA, Garcia PD, Klinger J, Kassam A;
XX Reinhard C, Randazzo F, Kennedy GC, Pot D, Lamson G, Drmanac R;
XX Ckenjakov R, Drmanac S, Dickson M, Labat I, Leshkowitz D;
XX Kila D, Garcia V, Jones LW, Strache-Train B;
XX
XX MPI: 2001-091805/10.
XX
XX Library of polynucleotides for diagnosing a cancerous state of a
XX mammalian cell and detecting cancer, particularly of the colon or
XX prostate, comprises 3351 human polynucleotide sequences -
XX

CC The invention relates to a single exon nucleic acid probe for

CC probes are useful for predicting, diagnosing, grading, staging, monitoring and prognosing diseases of the human heart and vascular system.

CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and

CC congenital heart disease.

CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/publised_sequences.
CC

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 52.4%; Score 58.2; DB 22; Length 570;

Best Local Similarity 79.3%; Pred. No. 8.2e-10;

Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

OY 24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAACTCTGGGCT 83

DB 566 GAATGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

OY 84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110

DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

ID AKK11985 standard; DNA: 570 BP.

XX AKK11985;

DT 05-NOV-2001 (first entry)

DE Human brain expressed single exon probe SEQ ID NO: 11976.

XX Human; brain expressed exon; gene expression analysis; probe;

KM microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;

KM epilepsy; cancer; ss.

OS Homo sapiens.

XX MO200157275-A2.

PN 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US00667.

PF 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

PA (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-483446/52.

XX Single exon nucleic acid probes for analyzing gene expression in human

PT brains -

PS Example 4; SEQ ID NO: 11976; 650bp + Sequence Listing; English.

XX The present invention provides a number of single exon nucleic acid

XX probes which are derived from genomic sequences expressed in the human

XX brain. They can be used to measure gene expression in brain cell samples,

XX CC which may enable the diagnosis and improved treatment of nervous system

XX diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,

XX epilepsy and cancers. The present sequence is one of the probes of the

XX invention.

Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

OY 24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAACTCTGGGCT 83

DB 566 GAATGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

OY 84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110

DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

ID AKK37688 standard; DNA: 570 BP.

XX AKK37688;

DT 06-NOV-2001 (first entry)

DE Human bone marrow expressed single exon probe SEQ ID NO: 12245.

XX Human; bone marrow expressed exon; gene expression analysis; probe;

KM microarray; cancer; leukaemia; lymphoma; myeloma; ss.

OS Homo sapiens.

XX MO200157276-A2.

PN 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US00668.

PF 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

PA (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-488900/53.

XX Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human bone marrow -

PS Example 4; SEQ ID NO: 12245; 658bp + Sequence Listing; English.

XX The present invention provides a number of single exon nucleic acid

XX probes which are derived from genomic sequences expressed in the human

XX bone marrow. They can be used to measure gene expression in bone marrow

XX samples, which may enable the improved diagnosis and treatment of cancers

XX CC such as lymphoma, leukaemia and myeloma. The present sequence is one of

XX the probes of the invention.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 52.4%; Score 58.2; DB 22; Length 570;

Best Local Similarity 79.3%; Pred. No. 8.2e-10; Mismatches 18; Indels 0; Gaps 0;

OY 24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAACTCTGGGCT 83

DB 566 GAATGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

OY 84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110

DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

RESULT 8
AA118447/c
ID AA118447 standard; DNA: 570 BP.
XX
XX
AC AA118447;
XX
XX
DT 12-OCT-2001 (first entry)
XX
XX
DE Probe #8380 for gene expression analysis in human cervical cell sample.
XX
XX
KW Probe: human; microarray; gene expression; cervical epithelial cell;
KW cervical cancer; ss.
XX
XX
OS Homo sapiens.
XX
XX
PN MO200157278-A2.
XX
XX
PD 09-AUG-2001.
XX
XX
30-JAN-2001; 2001MO-US00670.
XX
XX
04-FEB-2000; 2000US-0180312.
XX
XX
26-MAY-2000; 2000US-0207456.
XX
XX
30-JUN-2000; 2000US-0608408.
XX
XX
03-AUG-2000; 2000US-0632366.
XX
XX
21-SEP-2000; 2000US-0234687.
XX
XX
27-SEP-2000; 2000US-0236359.
XX
XX
04-OCT-2000; 2000GB-0024263.
XX
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX
DR WPI; 2001-488901/53.
XX
XX
PT Human genome-derived single exon nucleic acid probes useful for
XX
XX
PT analyzing gene expression in human cervical epithelial cells -
XX
XX
PS Claim 25; SEQ ID No 8380; 487bp; English.
XX
XX
CC The present invention relates to human single exon nucleic acid probes
XX
XX
CC (SENPs). The present sequence is one such probe. The SENPs are derived
XX
XX
CC from human HeLa cells. The SENPs can be used to produce a single exon
XX
XX
CC microarray, which can be used for measuring human gene expression in a
XX
XX
CC sample derived from human cervical epithelial cells. By measuring gene
XX
XX
CC expression, the probes are therefore useful in grading and/or staging
XX
XX
CC of diseases of the cervix, notably cervical cancer.
XX
XX
Note: The sequence data for this patent did not form part of the printed
XX
XX
specification, but was obtained in electronic format directly from WIPO
XX
XX
at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
XX
Query Match 52.4%; Score 58.2; DB 22; Length 570;
XX
XX
Best Local Similarity 79.3%; Pred. No. 8.2e-10;
XX
XX
Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
XX
XX
OY 24 GGATCTTTTTCATCTTTCAGAGGACTTCTGGGCGGAGATATATAAATCTGGGCT 83
XX
XX
DB 566 GAATGATCTCTCTGCTGCTTGGGATTCCTGGGCTGGAGTATATAAATTCCTGGGCT 507
XX
XX
OY 84 CTGTGTGTGCTGAGTGCTGCTTAC 110
XX
XX
DB 506 TTGTGTGTGCTGAGTGCGCTCTGC 480
XX
XX
RESULT 9
AA143563/c
ID AA143563 standard; DNA: 570 BP.
XX
XX
AC AA143563;
XX
XX
DT 17-OCT-2001 (first entry)

XX
XX
DE Probe #12249 used to measure gene expression in human placenta sample.
XX
XX
KW Probe: microarray; human; placenta; antenatal diagnosis;
KW genetic disorder; ss.
XX
XX
OS Homo sapiens.
XX
XX
PN MO200157272-A2.
XX
XX
PD 09-AUG-2001.
XX
XX
30-JAN-2001; 2001MO-US00663.
XX
XX
04-FEB-2000; 2000US-0180312.
XX
XX
26-MAY-2000; 2000US-0207456.
XX
XX
30-JUN-2000; 2000US-0608408.
XX
XX
03-AUG-2000; 2000US-0632366.
XX
XX
21-SEP-2000; 2000US-0234687.
XX
XX
27-SEP-2000; 2000US-0236359.
XX
XX
04-OCT-2000; 2000GB-0024263.
XX
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX
DR WPI; 2001-48897/53.
XX
XX
PT Human genome-derived single exon nucleic acid probes useful for
XX
XX
PT analyzing gene expression in human placenta -
XX
XX
PS Claim 25; SEQ ID No 12249; 654bp; English.
XX
XX
CC The present invention relates to single exon nucleic acid probes (SENPs).
XX
XX
CC The present sequence is one such probe. The probes are useful for
XX
XX
CC producing a microarray for predicting, measuring and displaying gene
XX
XX
CC expression in samples derived from human placenta. The probes are useful
XX
XX
CC for antenatal diagnosis of human genetic disorders.
XX
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
XX
Query Match 52.4%; Score 58.2; DB 22; Length 570;
XX
XX
Best Local Similarity 79.3%; Pred. No. 8.2e-10;
XX
XX
Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
XX
XX
OY 24 GGATCTTTTTCATCTTTCAGAGGACTTCTGGGCGGAGATATATAAATCTGGGCT 83
XX
XX
DB 566 GAATGATCTCTCTGCTGCTTGGGATTCCTGGGCTGGAGTATATAAATTCCTGGGCT 507
XX
XX
OY 84 CTGTGTGTGCTGAGTGCTGCTTAC 110
XX
XX
DB 506 TTGTGTGTGCTGAGTGCGCTCTGC 480
XX
XX
RESULT 10
ABS11680/c
ID ABS11680 standard; DNA: 570 BP.
XX
XX
AC ABS11680;
XX
XX
DT 19-AUG-2002 (first entry)
XX
XX
DE Human genome-derived single exon probe from lung SEQ ID No 11671.
XX
XX
KW Human: ds; single exon probe; asthma; lung cancer; COPD; ILD;
KW chronic obstructive pulmonary disease; interstitial lung disease;
KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karsenger syndrome;
KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
KW primary ciliary dyskinesia; pulmonary hypertension;
KW hyaline membrane disease.

XX Homo sapiens.
 OS
 XX WO200186003-A2.
 PN
 XX 15-NOV-2001.
 PD
 XX
 XX 30-JAN-2001; 2001WO-US00665.
 PF
 XX
 XX 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA
 XX Penn SC, Hanzel DK, Chen W, Rank DR;
 XX
 XX WPI: 2002-114183/15.
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 XX Claim 1: SEQ ID No 11671; 634pp; English.
 XX
 XX The invention relates to a spatially-addressable set of single exon
 XX nucleic acid probes for measuring gene expression in a sample derived
 XX from human lung comprising single exon nucleic acid probes having one of
 XX 12614 nucleic acid sequences mentioned in the specification, or their
 XX complements or the 12387 open reading frames derived from the 12614
 XX probes. Also included are a microarray comprising the novel set of
 XX probes; the novel set of probes which hybridize at high stringency to a
 XX nucleic acid expressed in the human lung; measuring gene expression in a
 XX sample derived from human lung, comprising (a) contacting the array with
 XX a collection of detectably labeled nucleic acids derived from human lung
 XX mRNA, and (b) measuring the label detectably bound to each probe of
 XX the array; identifying exons in a eukaryotic genome, comprising
 XX (a) algorithmically predicting at least one exon from genomic sequences
 XX of the eukaryote; and (b) detecting specific hybridisation of detectably
 XX labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 XX having a fragment identical to the predicted exon, the probe is included
 XX in the above mentioned microarray; assigning exons to a single gene,
 XX comprising (a) identifying exons from genomic sequence by the method
 XX above and (b) measuring the expression of each of the exons in several
 XX tissues and/or cell types using hybridisation to a single exon
 XX microarrays having a probe with the exon, where a common pattern of
 XX expression of the exons in the tissues and/or cell types indicates that
 XX the exons should be assigned to a single gene; a peptide comprising one
 XX of 12011 sequences, mentioned in the specification, or encoded by the
 XX probes/open reading frames (ORF). The probes are used for gene
 XX expression analysis, and for identifying exons in a gene, particularly
 XX using human lung derived mRNA and for the study of lung diseases
 XX such as asthma, lung cancer, chronic obstructive pulmonary disease
 XX (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 XX fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 XX Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 XX haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 XX pulmonary alveolar proteinosis, Karagenen syndrome, fibrocystic
 XX pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 XX and hyaline membrane disease. The present sequence is a single exon
 XX probe of the invention.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 CC
 XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
 Query Match 52.4%; Score 58.2; DB 24; Length 570;
 Best Local Similarity 79.3%; Pred. No. 8.2e-10;

Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
 QY 24 GCATTCCTTTTTCATCTTTGACGAGACTTCTGGGCGGAGATGTAACCTCTGGTCT 83
 Db 566 GAATGATCTCCCTGCTGCTGGATCTCTGGCTGAGATGTAACCTCTGGTCT 507
 QY 84 CTGTGTGTGCTGAGTGGCTGCTTAC 110
 Db 506 TTCTGTGTGCTGAGTGGCTGCTTCTGC 480
 RESULT 11
 ABR83461/C
 ID ABR83461 standard; cDNA; 128600 BP.
 XX
 XX ABR83461;
 AC
 XX
 XX 14-AUG-2002 (first entry)
 DT
 XX
 XX Human cDNA differentially expressed in granulocytic cells #32.
 DE
 XX
 XX Human; ss; granulocytic cell; DNA chip; bacterial infection;
 KW viral infection; parasitic infection; protozoal infection;
 KW fungal infection; sterile inflammatory disease; porriasis;
 KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
 KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.
 KW
 XX Homo sapiens.
 OS
 XX WO200228999-A2.
 PN
 XX 11-APR-2002.
 PD
 XX
 XX 03-OCT-2001; 2001WO-US30821.
 PF
 XX
 XX 03-OCT-2000; 2000US-237189P.
 PR
 XX
 XX (GENE-) GENE LOGIC INC.
 PA
 XX Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
 PI
 XX
 XX WPI: 2002-435328/46.
 PT
 PT Detecting granulocyte activation by detecting differential expression
 PT of genes associated with granulocyte activation, which serves as
 PT diagnostic markers that is useful for monitoring disease states and
 PT drug toxicity -
 XX
 XX Claim 1: SEQ ID No 32; 114pp; English.
 PS
 XX
 XX The invention relates to detecting (M1) granulocyte (GC) activation
 XX (GCA), by detecting the level of expression of gene(s) (Gs) identified by
 XX DNA chip analysis as given in the specification, and comparing
 XX the expression level to an expression level in an unactivated
 XX GC, where differential expression of Gs is indicative of GCA.
 XX Also included are modulating (M2) GA by contacting GC with an agent
 XX that alters the expression of at least one gene in Gs; (2) screening (M3)
 XX for an agent capable of modulating GCA or an inflammation (especially
 XX chronic) in a tissue, an allergic response in a subject, exposure of a
 XX subject to a pathogen or sterile inflammatory disease using the
 XX gene expression profile; (3) detecting (M4) an inflammation (especially
 XX chronic) in a tissue, an allergic response in a subject, exposure of a
 XX subject to a pathogen or sterile inflammatory disease, by detecting the
 XX level of expression in a sample of the tissue of gene(s) from Gs, where
 XX the level of expression of the gene is indicative of inflammation;
 XX (4) treating (M5) an inflammation (especially chronic) or in a tissue,
 XX an allergic response in a subject, exposure of a subject to a pathogen
 XX or sterile inflammatory disease, by contacting a tissue having
 XX inflammation with an agent that modulates the expression of gene(s)
 XX from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for

GenCore version 5.1.4-p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 01:47:50 ; Search time 1042 Seconds
(without alignments)
1725.239 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgacatcttgcctt.....gcttgagtgctgctact 111

Scoring table: IDENTITY_NUC
Gapop 10.0 ; Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

1 number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08
Maximum Match 1008

Listing first 45 summaries

Database :

BST:*

- 1: em_estba:*
- 2: em_esthum:*
- 3: em_estlin:*
- 4: em_estlm:*
- 5: em_estov:*
- 6: em_estpl:*
- 7: em_estro:*
- 8: em_hic:*
- 9: gb_est1:*
- 10: gb_est2:*
- 11: gb_hic:*
- 12: gb_est3:*
- 13: gb_est4:*
- 14: gb_est5:*
- 15: em_estlum:*
- 16: em_estom:*
- 17: gb_gss:*
- 18: em_gss_hum:*
- 19: em_gss_inv:*
- 20: em_gss_pln:*
- 21: em_gss_vrt:*
- 22: em_gss_fun:*
- 23: em_gss_mam:*
- 24: em_gss_mus:*
- 25: em_gss_other:*
- 26: em_gss_pro:*
- 27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	470 17	AQ770688	AQ770688 HS_5368_B
2	99.4	89.5	525 17	AQ165256	AQ165256 HS_3025_B
3	63.4	57.1	519 17	AQ186743	AQ186743 HS_3109_B
4	63	56.8	416 17	AQ182486	AQ182486 HS_3077_A
5	63	56.8	867 12	BG675164	BG675164 602621440
6	61.4	55.3	516 17	AQ457001	AQ457001 HS_5151_A

C	7	58.8	53.0	529	17	AQ881246	HS_5137_B
C	8	58.4	52.6	597	17	AQ506984	RPCT-11-3
C	9	58.2	52.4	416	17	B75884	RPCT11-12L1
C	10	58.2	52.4	530	17	AQ193128	HS_3060_B
C	11	57.8	52.1	448	17	AQ702903	HS_5452_B
C	12	57.8	52.1	769	17	AQ699390	HS_5234_A
C	13	57.6	51.9	413	17	AQ266658	RPCT11-73
C	14	57.2	51.5	453	17	AQ442274	HS_5137_A
C	15	57.2	51.5	635	17	AQ390599	HS_5137_A
C	16	56.8	51.2	519	17	AQ139984	HS_3106_A
C	17	56.6	51.0	491	17	AQ186162	HS_3077_A
C	18	56.6	51.0	628	17	AQ237815	RPCT11-70
C	19	56.6	51.0	864	17	AQ739814	HS_5505_A
C	20	56.6	50.5	546	17	AQ637256	RPCT-11-4
C	21	56	50.1	360	17	AQ207172	HS_3239_B
C	22	55.6	50.1	513	17	AQ455447	HS_5153_A
C	23	55.6	50.1	541	17	AQ765592	HS_5348_B
C	24	55.6	50.1	546	17	AQ541696	RPCT-11-3
C	25	55.6	50.1	546	17	AQ427698	CITBI-E1-
C	26	55.6	50.1	732	17	AQ506035	RPCT-11-3
C	27	55.6	50.1	580	17	AQ532835	RPCT-11-3
C	28	55.4	49.9	695	17	AG179297	Pan trogl
C	29	55.2	49.7	423	17	AQ564722	HS_5361_A
C	30	55	49.5	482	17	AQ320567	RPCT11-93
C	31	55	49.5	549	17	AQ40868	RPCT-11-3
C	32	55	49.5	551	17	AQ569689	HS_5333_B
C	33	55	49.5	563	17	AQ420187	RPCT-11-1
C	34	55	49.5	580	17	AQ161619	RPCT11-10
C	35	55	49.5	634	17	AG160901	Pan trogl
C	36	55	49.5	659	17	AG151043	Pan trogl
C	37	55	49.5	723	17	AQ386439	RPCT11-15
C	38	55	49.4	292	17	AQ508480	RPCT-11-2
C	39	54.8	49.4	483	17	AQ668395	HS_5414_A
C	40	54.6	49.2	653	17	AG143347	Pan trogl
C	41	54.6	49.2	683	17	AG091225	Pan trogl
C	42	54.6	49.2	503	17	AQ155611	HS_3124_A
C	43	54.4	49.0	708	17	AG092466	Pan trogl
C	44	54.4	49.0	672	17	AG051939	Pan trogl
C	45	54.2	48.8	17			

ALIGNMENTS

RESULT 1
LOCUS AQ770688
DEFINITION HS_5368_B2.C08.SP6E RPCT-11 Human Male BAC library Homo sapiens genomic clone Plate-944 Col-16 Row-F, DNA sequence.
ACCESSION AQ770688
VERSION AQ770688.1 GI:5648804
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 470)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
CONTACT: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCT-11. For BAC library availability, please contact Pieter de Jong

(piater@ejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering/bac.htm>) or from Research Genetics (<http://www.htsc.washington.edu>). BAC end Web Server: <http://www.htsc.washington.edu>

Plate: 944 Row: F Column: 16
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 470.

FEATURES

source

1..470

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone.lib="Plate=944 Col=16 Row=F"

/sex="male"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"

Query Match 100.0%; Score 111; DB 17; Length 470;
Best Local Similarity 100.0%; Pred. No. 2.3e-23;
Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGGACTTGGGCGG 60
|||||
Db 103 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGGACTTGGGCGG 162
|||||
Y 61 GAGTATGTAACACTCTGGTCTGTGTGTGCTGAGTGAGTGGCTGCTACT 111
|||||
Db 163 GAGTATGTAACACTCTGGTCTGTGTGTGCTGAGTGAGTGGCTGCTACT 213
|||||

RESULT 2

LOCUS

AQ165256 525 bp DNA linear GSS 16-OCT-1998

DEFINITION

HS.3025.B2.G06.T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3025 Col=12 Row=N, DNA sequence.

ACCESSION

AQ165256

VERSION

GSS.

KEYWORDS

human.

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

1 (bases 1 to 525)

Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and Hood, L.

Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

99380589

Contact: Mahairas GG, Wallace JC, Hood L

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 3025 Row: N Column: 12

Class: BAC ends

High quality sequence stop: 525.

Location/Qualifiers

1..525

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone.lib="Plate=3025 Col=12 Row=N"

/clone.lib="CIT Approved Human Genomic Sperm Library D"

FEATURES

source

/sex="male"
/note="Organ: sperm; Vector: pBelOBAC11; BAC clones in E-Coli DH10B"

Query Match 89.5%; Score 99.4; DB 17; Length 525;
Best Local Similarity 92.8%; Pred. No. 7.6e-20;
Matches 103; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

Y 1 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGGACTTGGGCGG 60
|||||
Db 66 ATGGGTGATCTTTTGGCTTGCAGATCTTTTCATCTTTGCGAGGACTTGGGCGG 125
|||||
Y 61 GAGTATGTAACACTCTGGTCTGTGTGTGCTGAGTGAGTGGCTGCTACT 111
|||||
Db 126 GAGTATGTAACACTCTGGTCTGTGTGTGCTGAGTGAGTGGCTGCTACT 176
|||||

RESULT 3

LOCUS

AQ16743 519 bp DNA linear GSS 01-NOV-1998

DEFINITION

HS.3109.B1.A06.T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3109 Col=11 Row=B, DNA sequence.

ACCESSION

AQ16743

VERSION

GSS.

KEYWORDS

human.

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

1 (bases 1 to 519)

Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and Hood, L.

Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

99380589

Contact: Mahairas GG, Wallace JC, Hood L

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 3109 Row: B Column: 11

Class: BAC ends

High quality sequence stop: 519.

Location/Qualifiers

1..519

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone.lib="Plate=3109 Col=11 Row=B"

/clone.lib="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBelOBAC11; BAC clones in E-Coli DH10B"

BASE COUNT

144 a 155 c 122 g 93 t 5 others

ORIGIN

Query Match

Best Local Similarity

Matches 73; Conservative

0; Mismatches 16; Indels 0; Gaps 0;

Y 23 AGGATTCCTTTTCATCTTTGCGAGGACTTCTGGGCGGAGTATGTAACACTCTGGGTC 82

|||||

Db 398 AGGATTCCTTTTCATCTTTGCGAGGACTTCTGGGCGGAGTATGTAACACTCTGGGTC 339

Y 83 TCTGTGTGCTGAGTGGCTGCTACT 111

|||||

Db 338 TCTGTGTGCTGAGTGGCTGCTACT 310

[illegible]

JOURNAL	Unpublished (1999)
COMMENT	Contact: Robert Strausberg, Ph.D. Email: cgapbs-remail.nih.gov Tissue Procurement: James Cleaver, M.D. cDNA Library Preparation: Life Technologies, Inc. CDNA library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Incyte genomics, Inc. Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: LLAM10617 row: c column: 09 High quality sequence start: 5 High quality sequence stop: 857.
FEATURES	Location/Qualifiers
SOURCE	1..867 /organism="Homo sapiens" /db_xref="taxon:9606" /clone="IMAGE:4755296" /clone_lib="NCI CGAP_Skn3" /lab_host="DH10B (TI phage-resistant)" /note="Organ: skin; Vector: pGMV-SPOrtG; Site_1: NotI; Site_2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.5kb. Library constructed by Life Technologies. Note: This is a NCI-Cgap Library."
BASE COUNT	285 A 208 C 217 G 157 T
ORIGIN	
Query Match	56.8%; Score 63; DB 12; Length 867;
Best Local Similarity	73.0% ; Pred. No. 9.2e+09;
Matches	81; Conservative 0; Mismatches 30; Indels 0; Gaps 0.
OY	1 ATGGCGATCTTTCCCTTCAGCATCTCTTGTACATTCTTGCAAGCATTCTGGGCCG 60
Dd	636 ATTGATGACCCTCCCCCTTAGACTAGTGTAAGTGAACCTTGCCAGGAGATCTGGTGC 577
OY	61 GAGTATGTAACAATCCTGGGCTCTCTGTGTGTGCGCTGAAAGTGCCTCTACT 111
Dd	576 GGATATATAAATCTTGAGTCTGTGTGATCAATCACAGAATGGCTGCTGCT 526
RESULT 6	
AQ457001 LOCUS	Hs_515L.AL.G09.T7A.RPCT-11 Human Male BAC Library Homo sapiens genomic clone Plate=727 Col=17 Row=M, DNA sequence.
ACCESSION	AQ457001
VERSION	AQ457001.1 GI:4635878
KEYWORDS	GSS.
SOURCE ORGANISM	human. Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE AUTHORS	1 (bases 1 to 516) Mahairas,G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L. Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
TITLE	* Contact: Mahairas GG, Wallace JC, Hood L
JOURNAL MEDLINE COMMENT	High Throughput Sequencing Center University of Washington 401 Queen Anne Avenue North, Seattle, WA 98109, USA Tel.: (206) 616-3618 Fax: (206) 616-3887 Email: jwallace@u.washington.edu Clones are derived from the human BAC library RPCT-11. For BAC library availability, please contact Pieter de Jong (pjeterdejong.med.buffalo.edu). Clones may be purchased from BACPac Resources (http://bacpac.med.buffalo.edu/ordering/bac.htm) or from Resear h Genetics (inforesgen.com). BAC end Web Server: http://www.htsc.washington.edu

FEATURES
source
Location/Qualifiers
1. 413
/organism="Homo sapiens"
/db_xref="GDB:7527878"
/db_xref="taxon:9606"
/clone_id="RPCI-11-73015"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: PBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC Library"

BASE COUNT
75 a 108 c 118 g 112 t

ORIGIN
Query Match 51.5%; Score 57.6; DB 17; Length 413;
Best Local Similarity 78.4%; Pred. No. 3.5e-07;
Matches 69; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

OY 24 GGATCTTTTCATCTTGCAGGACTTTCGGGCGGAGTATGTAACCTCGGCTC 83
104 GGATGATCTCCCATCTTGCAAGAACCTGCGCTGAGTATGCAAACTCCTGGCTC 163

OY 84 CTGTGTGCTGCTGAGTGTGCTCTCTACT 111
164 CTGTGTGCTGCTGAGTGTGCTCTCTGCT 191

RESULT 14
LOCUS A0442274 410 bp DNA linear GSS 31-MAR-1999
DEFINITION HS.5137.A1.F12.SPE6 RPCI-11 Human Male BAC Library Homo sapiens
ACCESSION A0442274
VERSION A0442274.1 GI:4553613
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 410)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (inforesgen.com). BAC end web Server:
http://www.htsc.washington.edu
Plate: 713 row: K column: 23
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 410.
Location/Qualifiers
1. 410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_id="plate=713 Col=23 Row=K"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: PBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor

FEATURES
source
Location/Qualifiers
1. 410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_id="plate=713 Col=23 Row=K"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: PBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor

BASE COUNT
80 a 111 c 111 g 107 t 1 others

ORIGIN
Query Match 51.5%; Score 57.2; DB 17; Length 410;
Best Local Similarity 79.1%; Pred. No. 4.6e-07;
Matches 68; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

OY 25 GATCTTTTCATCTTGCAGGACTTTCGGGCGGAGTATGTAACCTCGGCTC 84
111 GATGCTTCTCTGCTGCTGCTGATCCAGGCGCAGAGTATGCAAACTCCTGGCTC 170

OY 85 TGTGTGCTGCTGAGTGTGCTCTCTAC 110
171 TGTGTGCTGCTGAGTGTGCTCTCTGC 196

RESULT 15
LOCUS A0437684 453 bp DNA linear GSS 31-MAR-1999
DEFINITION HS.5137.A2.H06.SPE6 RPCI-11 Human Male BAC Library Homo sapiens
ACCESSION A0437684
VERSION A0437684.1 GI:4549023
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 453)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (inforesgen.com). BAC end web Server:
http://www.htsc.washington.edu
Plate: 713 row: O column: 12
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 453.
Location/Qualifiers
1. 453
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_id="plate=713 Col=12 Row=O"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: PBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
PBACe3.6 vector at EcoRI sites"

BASE COUNT
84 a 127 c 117 g 124 t 1 others

ORIGIN
Query Match 51.5%; Score 57.2; DB 17; Length 453;
Best Local Similarity 79.1%; Pred. No. 4.7e-07;

	Matches	68;	Conservative	0;	Mismatches	18;	Indels	0;	Gaps	0;
QY	25	GATTCTTTTTCATCTTTGAGGACTTCTGGGCGCGAGATATGTA	AACTCCTGGGCTC	84						
Db	114	GATGGTTCTTCTGCTGCTGATGCCAGGCGCAGAGTATGCA	AAATTCCTGGGCTC	173						
QY	85	TGTGTGTGCTGAGTGGCTGCTTAC	110							
Db	174	TGTGTGTGCTGAGTGGCTGCTTGC	199							

Search completed: April 25, 2003, 02:28:38
Job time : 1045 secs

GenCore version 5.1.4-P5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 01:55:25 ; Search time 39 Seconds
(without alignments)
872.849 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111

Sequence: 1 atgggtgacatttgcctt.....gcctgagtgctgcttact 111

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 441362 seqs, 15338381 residues

1 number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.*

1: /cgn2_6/ptodata/1/ina/5A.COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B.COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A.COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B.COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCUTS.COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfile1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	27.6	24.9	80246	4	US-09-078-294-4
2	27.6	24.9	80595	4	US-09-078-294-3
3	26.2	23.6	175	1	US-08-222-177A-4
4	26.2	23.6	1146	2	US-08-666-367B-4
5	26.2	23.6	1146	4	US-09-143-438-4
6	26.2	23.6	13104	4	US-08-961-527-34
7	26.2	23.4	430	4	US-09-397-787-254
8	25.4	22.9	11492	4	US-08-991-840A-1
9	24.8	22.3	1371	2	US-08-428-713-1
10	24.8	22.3	1371	3	US-08-904-179-1
11	24.8	22.3	1374	2	US-08-428-713-9
12	24.8	22.3	1374	3	US-08-904-179-9
13	24.6	22.2	882	4	US-09-556-877-136
14	24.6	22.2	882	4	US-09-620-412C-136
15	24.6	22.2	2407	4	US-09-370-807-7
16	24.6	22.2	2407	4	US-09-921-259-7
17	24.6	22.2	2511	4	US-09-422-869-19
18	24.4	22.0	361	4	US-09-385-982-26
19	24.4	22.0	1506	4	US-08-206-790A-22
20	24.4	22.0	1506	5	PCT-US95-02843-22
21	24.2	21.8	2347	1	US-08-145-681-3
22	24.2	21.8	2347	1	US-08-453-703-3
23	24.2	21.8	2347	2	US-08-456-106-3
24	24.2	21.8	2347	3	US-08-456-108-3
25	24.2	21.8	2347	4	US-09-265-577-3
26	24.2	21.6	528	4	US-09-134-001C-2094
27	24.2	21.6	4088	2	US-08-317-310A-1

28	24	21.6	4088	5	PCT-US95-13041-1	Sequence 1, Appli
29	24	21.6	19446	4	US-08-961-527-51	Sequence 51, Appl
30	23.8	21.4	3831	4	US-08-961-527-291	Sequence 291, App
31	23.8	21.4	7542	4	US-09-734-030-3	Sequence 3, Appli
32	23.8	21.4	13158	2	US-08-687-080-105	Sequence 105, App
33	23.8	21.4	35100	1	US-08-306-691B-19	Sequence 19, Appl
34	23.8	21.4	35100	3	PCT-US93-06251-19	Sequence 19, Appl
35	23.8	21.4	87350	3	US-08-781-891-79	Sequence 79, Appl
36	23.8	21.4	87543	4	US-09-791-211-3	Sequence 3, Appli
37	23.6	21.3	15567	4	US-09-627-376-3	Sequence 3, Appli
38	23.4	21.1	560	3	US-09-059-369-18	Sequence 18, Appl
39	23.4	21.1	598	4	US-08-998-416-1076	Sequence 1076, Ap
40	23.4	21.1	694	4	US-08-998-416-710	Sequence 710, App
41	23.4	21.1	1477	3	US-09-123-030-9	Sequence 9, Appli
42	23.4	21.1	2090	3	US-09-059-369-1	Sequence 1, Appli
43	23.4	21.1	3093	1	US-08-252-966B-19	Sequence 19, Appli
44	23.4	21.1	5253	2	US-08-290-731C-3	Sequence 3, Appli
45	23.2	20.9	401	4	US-09-221-296-40	Sequence 40, Appli

ALIGNMENTS

```
RESULT 1
US-09-078-294-4
; Sequence 4, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078, 294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 4
; LENGTH: 80246
; TYPE: DNA
; ORGANISM: Nucleotide sequence of NC-contlig
US-09-078-294-4

Query Match      24.9%; Score 27.6; DB 4; Length 80246;
Best Local Similarity 58.5%; Pred. No. 5.3;
Matches 48; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

QY 23 AGCATTTCTTTTATCTTTCAGGAGCTTGGGCCGAGATGTAACACTCTGGGTC 82
    || |||| |||| |||| |||| |||| |||| |||| |||| |||| ||||
Db 6333 AGTTTCTGTGTCAACTGTGACGTGGCCATGGCATGTGTCAGATATGTAATTAAACAGTATT 6392
QY 83 TCTGTGTGCTGCTGAGTGGCTG 104
    |||| |||| |||| |||| |||| |||| |||| |||| |||| ||||
Db 6393 TCTGTGTGTTTCTGTGAGGGTG 6414

RESULT 2
US-09-078-294-3
; Sequence 3, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078, 294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 3
; LENGTH: 80595
; TYPE: DNA
```



```

OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/666,367B
FILING DATE: August 19, 1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Warren M. Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-8850
TELEFAX:
TELEX:
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1146
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA
ORIGINAL SOURCE:
ORGANISM: mouse
FEATURE:
OTHER INFORMATION: 1-1128 sialyltransferase in soluble
US-08-666-367B-4

Query Match      23.6%; Score 26.2; DB 2; Length 1146;
Best Local Similarity 63.5%; Pred. No. 3.5;
Matches 40; Conservative 0; Mismatches 23; Indels 0; Gaps 0;

QY 19 TTGCAGATCTTTTCATCTTGCAGGACTCTGCGCGGAGTATGTAACCTCTG 78
    || || || || || || || || || || || || || || || || || || ||
DB 840 TTCTGATGATGTCCTCATGTCCTCATGCGCTTGAGGATGTAAGGCTG 781
    || || || || || || || || || || || || || || || || || || ||

QY 79 GGT 81
DB 780 GCT 778

RESULT 5
US-09-143-438-4/c
Sequence 4, Application US/09143438
Patent No. 6218161
GENERAL INFORMATION:
APPLICANT: Shuichi TSUI et al.
TITLE OF INVENTION: NOVEL SUGAR-CHAIN SYNTHETASE AND PROCESS FOR
TITLE OF INVENTION: PRODUCING THE SAME
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: Wenderoth, Lind & Ponack, L.L.P.
STREET: 2033 K Street, N.W., #800
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20006
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 mb
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/143,438
FILING DATE: August 28, 1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/666,367
FILING DATE: August 19, 1996
ATTORNEY/AGENT INFORMATION:

```

```

NAME: Warren M. Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-721-8200
TELEFAX: 202-721-8250
TELEX:
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1146
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA
ORIGINAL SOURCE:
ORGANISM: mouse
FEATURE:
OTHER INFORMATION: 1-1128 sialyltransferase in soluble
US-09-143-438-4

Query Match      23.6%; Score 26.2; DB 4; Length 1146;
Best Local Similarity 63.5%; Pred. No. 3.5;
Matches 40; Conservative 0; Mismatches 23; Indels 0; Gaps 0;

QY 19 TTGCAGATCTTTTCATCTTGCAGGACTCTGCGCGGAGTATGTAACCTCTG 78
    || || || || || || || || || || || || || || || || || || ||
DB 840 TTCTGATGATGTCCTCATGTCCTCATGCGCTTGAGGATGTAAGGCTG 781
    || || || || || || || || || || || || || || || || || || ||

QY 79 GGT 81
DB 780 GCT 778

RESULT 6
US-08-961-527-34/c
Sequence 34, Application US/08961527
Patent No. 6420135
GENERAL INFORMATION:
APPLICANT: Charles Kunsch
TITLE OF INVENTION: Streptococcus pneumoniae Polynucleotides and Sequences
NUMBER OF SEQUENCES: 391
CORRESPONDENCE ADDRESS:
ADDRESSEE: Human Genome Sciences, Inc.
STREET: 9410 Key West Avenue
CITY: Rockville
STATE: Maryland
COUNTRY: USA
ZIP: 20850
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44mb storage
COMPUTER: HP Vectra 486/33
OPERATING SYSTEM: MSDOS version 6.2
SOFTWARE: ASCII text
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/961,527
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Brookes, A. Anders
REGISTRATION NUMBER: 36,373
REFERENCE/DOCKET NUMBER: PB340P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301) 309-8512
TELEFAX: (301) 309-8504
INFORMATION FOR SEQ ID NO: 34:
SEQUENCE CHARACTERISTICS:
LENGTH: 13104 base pairs
TYPE: nucleic acid
STRANDEDNESS: double

```

TOPOLOGY: Linear
US-08-961-527-34

Query Match 23.6%; Score 26.2; DB 4; Length 13104;
Best Local Similarity 67.3%; Pred. No. 8.6;
Matches 37; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

Db 9341 TCTATTGCTTGGGGGCTTTGGGCACTGTTTGAAGGAGTTTAAGCACGCT 9287

RESULT 7

US-09-397-787-254/C
Sequence 254; Application US/09397787
Patent No. 6468158
GENERAL INFORMATION:
APPLICANT: Benson, Darin R.
APPLICANT: Lodes, Michael J.
APPLICANT: Mitcham, Jennifer L.
APPLICANT: King, Gordon E.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
TITLE OF INVENTION: CANCER THERAPY AND DIAGNOSIS
FILE REFERENCE: 210121.466C2
CURRENT APPLICATION NUMBER: US/09/397.787
CURRENT FILING DATE: 1999-09-16
NUMBER OF SEQ ID NOS: 334
SOFTWARE: PastSeq for Windows Version 3.0
SEQ ID NO 254
LENGTH: 430
TYPE: DNA
ORGANISM: Homo sapien
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(430)
OTHER INFORMATION: n = A,T,C or G
US-09-397-787-254

Query Match 23.4%; Score 26; DB 4; Length 430;
Best Local Similarity 70.0%; Pred. No. 2.9;
Matches 35; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

Db 402 AGATATGAACCTCGGCTCTGTGTGTCGAGTGCTGCTACT 111
402 AGATATGAAGCTCCCAAGTCTCATGTCTTGTGGGGCTCTGCT 353

RESULT 8

US-991-840A-1/C
Sequence 1; Application US/08991840A
Patent No. 6261570

GENERAL INFORMATION:
APPLICANT: Michael D. Parker
APPLICANT: Jonathan F. Smith
APPLICANT: Bruce Crise
APPLICANT: Mark Steve Oberste
APPLICANT: Shannon Schmura
TITLE OF INVENTION: Live Attenuated Virus Vaccines for Eastern Equine Encephalitis
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: Charles H. Harris
STREET: USA MRC - MRC-JA
CITY: FORT DETRICK, FREDERICK
STATE: MARYLAND
COUNTRY: USA
ZIP: 21702-5012

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Macintosh 7.5
SOFTWARE: Microsoft Word 6.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/991.840A

FILING DATE: December 16, 1997
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: Provisional Application 60/047162,
FILING DATE: May 20, 1997
APPLICATION NUMBER: Provisional Application 60/053,652
FILING DATE: July 24, 1997

ATTORNEY/AGENT INFORMATION:
NAME: Charles H. Harris
REGISTRATION NUMBER: 34,616
REFERENCE/DOCKET NUMBER: 003/058/SAP RIID 96-01
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301) 619-2065
TELEFAX: (301) 619-5034
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 11492 base pairs
TYPE: Nucleic acid
STRANDEDNESS: Double

TOPOLOGY: Linear
FEATURE: OTHER INFORMATION: N at all occurrences is = unknown.
FEATURE: OTHER INFORMATION: K at all
FEATURE: occurrences is = G or T
US-08-991-840A-1

Query Match 22.9%; Score 25.4; DB 4; Length 11492;
Best Local Similarity 64.4%; Pred. No. 16;
Matches 38; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

Db 7692 TTTGGCTTGAGACACTTCTTCTTCTTGGGGGAGCCTGGCGGAGTTAGCTGA 7634

RESULT 9

US-08-428-713-1/C
Sequence 1; Application US/08428713
Patent No. 5866541

GENERAL INFORMATION:
APPLICANT: HOOK, Magnus
APPLICANT: LINDBERG, Kjell Martin
APPLICANT: LINDGREN, Per-Eric
APPLICANT: STENAS, Lars Christer
TITLE OF INVENTION: FIBRONECTIN BINDING PROTEIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Burns, Doane, Swecker & Mathis
STREET: P.O. Box 1404
CITY: Alexandria
STATE: Virginia
COUNTRY: United States
ZIP: 22313-1404

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/428.713
FILING DATE: 25-APR-1995
CLASSIFICATION: 514

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/125,222
FILING DATE: 23-SEP-1993
ATTORNEY/AGENT INFORMATION:
NAME: Rea, Teresa Stanek
REGISTRATION NUMBER: 30,427
REFERENCE/DOCKET NUMBER: 012885-074
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 836-6620
TELEFAX: (703) 836-2021
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:

Db 882 TTTTACAGCACTGCATTTGAATTCAGACACACAGCAATTAGACTGCTGCTTTAAACA 941
OY 90 GTGCCTGAGTGGCTGCTCT 108
| | | | | | | | | |
Db 942 TTCTCTGATTAAGTGTGT 960

Search completed: April 25, 2003, 02:29:59
Job time : 67 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 02:28:45 ; Search time 69 Seconds
(without alignments)
1750.468 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggttgatcttgcctt.....gcttgatgctgctact 111

Scoring table:
IDENTITY-NUC
Gapop 10.0 , Gapext 1.0

Searched: 709820 seqs, 544064369 residues

1 number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : Published Applications_NA:*
1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
2: /cgn2_6/ptodata/1/pubpna/PTCT_NEW_PUB.seq:*
3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
6: /cgn2_6/ptodata/1/pubpna/PTCTUS_PUBCOMB.seq:*
7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	58.2	52.4	570	US-09-864-761-9118	Sequence 9118, App
2	57.6	51.9	684973	US-09-263-959-1	Sequence 1, App11
3	52.8	47.6	428	US-09-918-993-8153	Sequence 8153, App
4	38.8	35.0	162	US-09-864-761-31393	Sequence 31393, A
5	38.8	35.0	519	US-09-864-761-14866	Sequence 14866, A
6	29.2	26.3	369	US-09-954-456-922	Sequence 922, App
7	29.2	26.3	1314	US-09-954-456-1538	Sequence 1538, App
8	29.2	26.3	1314	US-10-125-540-609	Sequence 609, App
9	29.2	26.3	1314	US-09-764-870-609	Sequence 609, App
10	28	25.2	342	US-10-125-540-118	Sequence 118, App
11	28	25.2	342	US-09-764-870-118	Sequence 118, App
12	27.2	24.5	412	US-09-783-590-1450	Sequence 1450, App
13	27.2	24.5	504	US-10-092-154-1665	Sequence 1665, App
14	27.2	24.5	504	US-10-092-154-1666	Sequence 1666, App
15	27.2	24.5	504	US-10-092-154-1667	Sequence 1667, App
16	27.2	24.5	504	US-09-764-847-1665	Sequence 1665, App
17	27.2	24.5	504	US-09-764-847-1666	Sequence 1666, App
18	27.2	24.5	504	US-09-764-847-1667	Sequence 1667, App
19	27	24.3	487	US-09-747-155-270	Sequence 270, App

20	26.6	24.0	6372	US-09-880-107-3948	Sequence 3948, App
21	26.4	23.8	385	US-09-770-791-90	Sequence 90, App1
22	26.4	23.8	1666	US-09-938-842A-966	Sequence 966, App
23	26.4	23.8	170834	US-09-835-232-7	Sequence 7, App11
24	26.2	23.6	26225	US-10-091-504-1276	Sequence 1276, App
25	26.2	23.6	26225	US-09-764-869-1276	Sequence 1276, App
26	26	23.4	430	US-09-876-889-254	Sequence 254, App
27	26	23.4	1548	US-09-925-302-211	Sequence 211, App
28	26	23.4	1620	US-09-954-456-325	Sequence 325, App
29	26	23.4	5641	US-10-015-219-1733	Sequence 1733, App
30	26	23.4	5709	US-10-015-219-1734	Sequence 1734, App
31	25.8	23.2	590	US-09-864-761-12014	Sequence 12014, A
32	25.8	23.2	1663	US-09-822-830A-248	Sequence 248, App
33	25.6	23.1	440	US-10-092-154-366	Sequence 366, App
34	25.6	23.1	440	US-09-764-847-366	Sequence 366, App
35	25.6	23.1	442	US-09-880-107-1917	Sequence 1917, App
36	25.6	23.1	1184	US-09-969-347-210	Sequence 210, App
37	25.6	23.1	32169	US-10-092-154-1963	Sequence 1963, App
38	25.6	23.1	32169	US-09-764-847-1963	Sequence 1963, App
39	25.4	22.9	11617	US-09-860-670-265	Sequence 265, App
40	25.2	22.7	390	US-09-770-791-11	Sequence 11, App1
41	25.2	22.7	455	US-09-960-352-9392	Sequence 9392, App
42	25.2	22.7	509	US-09-992-598-149	Sequence 149, App
43	25.2	22.7	509	US-09-989-735A-149	Sequence 149, App
44	25.2	22.7	509	US-09-989-735-149	Sequence 149, App
45	25.2	22.7	509	US-09-990-444-149	Sequence 149, App

ALIGNMENTS

RESULT 1
US-09-864-761-9118/c
Sequence 9118, Application US/09864761
Patient No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-06-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30

PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
SEQ ID NO 9118
LENGTH: 570
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000053.1
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.8
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
US-09-864-761-9118

Query Match 52.4%; Score 58.2; DB 10; Length 570;
Best Local Similarity 79.3%; Pred. No. 1.2e-10;
Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 24 GGATTCCTTTTCATCTTTCAGAGGACTTTCGGGCGGAGTATGTAATAACTCTGGGCT 83
DB 566 GAATGAGATCTCTCCCTGCTGGGATTCCTTGGGCTGAGATGTAATAATCTGGGCT 507
OY 84 CTGTGTGCTGCTGAGTGGCTGCTTAC 110
DB 506 TTGTGTGCTGCTGAGTGGCGCTCTGC 480

RESULT 2

US-09-263-959-1/c
Sequence 1, Application US/09263959
Patent No. US20020150891A1
GENERAL INFORMATION:
APPLICANT: Hood, Leroy E.
APPLICANT: Rowen, Lee
APPLICANT: Koop, Ben F.
TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH UTILIZE
NUMBER OF SEQUENCES: 1279
CORRESPONDENCE ADDRESS:
ADDRESSEE: Seed and Berry LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: US
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/263,959
FILING DATE: 05-MAR-1999
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 684973 base pairs
TYPE: nucleic acid
STRADEDNESS: single
TOPOLOGY: linear
US-09-263-959-1

Query Match 51.9%; Score 57.6; DB 10; Length 684973;
Best Local Similarity 78.4%; Pred. No. 1.5e-09;
Matches 69; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 24 GGATTCCTTTTCATCTTTCAGAGGACTTTCGGGCGGAGTATGTAATAACTCTGGGCT 83
DB 404732 GGACAGATTCTCCTGCTGGGAGTCTGAGCTGGAATGTAATAACTCTGGGCT 404673
QY 84 CTGTGTGCTGCTGAGTGGCTGCTTAC 111
DB 404672 CTGTGTGCTGCTGAGTGGCTGCTTAC 404645

RESULT 3

US-09-918-995-8153
Sequence 8153, Application US/09918995
Publication No. US20030073623A1
GENERAL INFORMATION:
APPLICANT: Hyseq, Inc.
TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED FROM VARIOUS CDNA LIBRARIES
FILE REFERENCE: 20411-756
CURRENT APPLICATION NUMBER: US/09/918,995
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US/09/235,076
PRIOR FILING DATE: 1999-01-20
NUMBER OF SEQ ID NOS: 38054
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 8153
LENGTH: 428
TYPE: DNA
ORGANISM: Homo sapiens
US-09-918-995-8153

Query Match 47.6%; Score 52.8; DB 9; Length 428;
Best Local Similarity 71.9%; Pred. No. 8.2e-09;
Matches 69; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

QY 15 TGCCCTGAGAGTCTTTTCATCTTTCAGAGGACTTTCGGGCGGAGTATGTAATAACT 74
DB 272 TGCCCTGAGAGTCTTTTCATCTTTCAGAGGACTTTCGGGCGGAGTATGTAATAACT 331
QY 75 CCGGCTGCTGCTGAGTGGCTGCTTAC 110
DB 332 CCGGCTGCTGAGTGGCTGCTTAC 367

RESULT 4

US-09-864-761-31393
Sequence 31393, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aecoliga-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366


```

: PRIOR FILLING DATE: 2000-08-03
: PRIOR APPLICATION NUMBER: GB 24263.6
: PRIOR FILLING DATE: 2000-10-04
: PRIOR APPLICATION NUMBER: US 60/236,359
: PRIOR FILLING DATE: 2000-09-27
: PRIOR APPLICATION NUMBER: PCT/US01/00666
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00667
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00664
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00669
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00665
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00668
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00663
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00662
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00661
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00670
: PRIOR FILLING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: US 60/234,687
: PRIOR FILLING DATE: 2000-09-21
: PRIOR APPLICATION NUMBER: US 09/608,408
: PRIOR FILLING DATE: 2000-06-30
: PRIOR APPLICATION NUMBER: US 09/774,203
: PRIOR FILLING DATE: 2001-01-29
: NUMBER OF SEQ ID NOS: 49117
: SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
: SEQ ID NO 31393
: LENGTH: 162
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: OTHER INFORMATION: MAP TO AC017089.2
: OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
: OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
: OTHER INFORMATION: SWISSPROT HIT: P98161, EVALUE 3.00e+00
: OTHER INFORMATION: EST HUMAN HIT: A1792950.1, EVALUE 6.00e-07
: OTHER INFORMATION: NT HIT: AL163210.2, EVALUE 4.00e-04
US-09-864-761-31393
Query Match 35.0%; Score 38.8; DB 10; Length 162;
Best Local Similarity 70.3%; Pred. No. 0.00043;
Matches 52; Conservative 0; Mismatches 22; Indels 0; Gaps 0;
●
OY 38 CTTTGACAGGACTTCTGGGGCCGAGATATGTAAACTCCTGGGTCCTGTGTGCTGCTTA 97
||||| 111 11 11111 11 11111 111 11 111111111
Db 52 CTTTCTTCGAGGACGAGGGCCAGATATCTAAAGCTCTGGAGTTTCAATGTGTGCTTA 111
||| 98 GTGGCTGCTCTACT 111
| 111111111
Db 112 GCACATGCTCTGCT 125
RESULT 5
US-09-864-761-14866
: Sequence 14866, Application US/09864761
: Patent No. US20020048763A1
: GENERAL INFORMATION:
: APPLICANT: Penn, Shatton G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
: TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
: FILE REFERENCE: Aeonica-X-1
: CURRENT APPLICATION NUMBER: US/09/864,761
: CURRENT FILING DATE: 2001-05-23

```

```

PRIORITY APPLICATION NUMBER: US 60/180,312
PRIORITY FILING DATE: 2000-02-04
PRIORITY APPLICATION NUMBER: US 60/207,456
PRIORITY FILING DATE: 2000-05-26
PRIORITY APPLICATION NUMBER: US 09/632,366
PRIORITY FILING DATE: 2000-08-03
PRIORITY APPLICATION NUMBER: GB 24263.6
PRIORITY FILING DATE: 2000-10-04
PRIORITY APPLICATION NUMBER: US 60/236,359
PRIORITY FILING DATE: 2000-09-27
PRIORITY APPLICATION NUMBER: PCT/US01/00666
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00667
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00664
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00669
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00665
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00668
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00663
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00662
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00661
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: PCT/US01/00670
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: US 60/234,687
PRIORITY FILING DATE: 2000-09-21
PRIORITY APPLICATION NUMBER: US 09/608,408
PRIORITY FILING DATE: 2000-06-30
PRIORITY APPLICATION NUMBER: US 09/774,203
PRIORITY FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
SEQ ID NO 14866
LENGTH: 519
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AC017089.2
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
US-09-864-761-14866

Query Match      35.0%   Score 38.8   DB 10;   Length 519;
Best Local Similarity 70.3%   Pred. No. 0.0006;
Matches 52; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

OY      38  CTTTGACAGGACTCTGGGCGGAGATGTAACATCTGCTCTGTGTGCGCTGA 97
        |||||  |||  ||  |||||  ||  |||||  |||  ||  ||  |||||  |||||
Db      301  CTTTCCCTCGAAGTCCAGAGCGCACGATATATCAAGCTGTGAGTTTCATGTCGCTGA 360
        |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

OY      98  GTGGCTGCTTACT 111
        |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db      361  GCAGATGCTGCT 374

RESULT 6
Sequence 922: Application US/09954456
Patent No. US20020115057A1
GENERAL INFORMATION:
APPLICANT: Young, Paul
TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using C
FILE REFERENCE: 689290-76
CURRENT APPLICATION NUMBER: US/09/954,456
PRIORITY FILING DATE: 2001-09-18
PRIORITY APPLICATION NUMBER: US/60/233,617

```

```

; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 922
; LENGTH: 369
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-922

Query Match          26.3%; Score 29.2; DB 10; Length 369;
Best Local Similarity 57.8%; Pred. No. 1.1;
Matches 52; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTGGCTTCGAGAGATCTTTTCATCTTTGACGAGGACTTCTGGGCGG 60
DQ 143 AGGCGTGAATAATTGATTGTTCAATATAGCTTCACGCTTAGCAATTAACCTAGTCCAA 84
DB 61 GAGTATGTAACACTCTGGGCTCTGTGTG 90
DB 83 GACAATATTGATTCCTAGTCTGTGTGG 54

RESULT 7
US-09-954-456-1538/C
; Sequence 1538, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents using Candi-
; TITLE OF INVENTION: Sets
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; CURRENT FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
```

```

; SEQ ID NO 1538
; LENGTH: 369
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-1538

Query Match          26.3%; Score 29.2; DB 10; Length 369;
Best Local Similarity 57.8%; Pred. No. 1.1;
Matches 52; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTGGCTTCGAGAGATCTTTTCATCTTTGACGAGGACTTCTGGGCGG 60
DQ 143 AGGCGTGAATAATTGATTGTTCAATATAGCTTCACGCTTAGCAATTAACCTAGTCCAA 84
DB 61 GAGTATGTAACACTCTGGGCTCTGTGTG 90
DB 83 GACAATATTGATTCCTAGTCTGTGTGG 54

RESULT 8
US-10-125-540-609
; Sequence 609, Application US/10125540
; Publication No. US20030059875A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214C1
; CURRENT APPLICATION NUMBER: US/10/125,540
; PRIOR FILING DATE: 2002-04-19
; PRIOR APPLICATION REMOVED - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 609
; LENGTH: 1314
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-125-540-609

Query Match          26.3%; Score 29.2; DB 9; Length 1314;
Best Local Similarity 56.1%; Pred. No. 1.6;
Matches 55; Conservative 0; Mismatches 43; Indels 0; Gaps 0;

QY 3 GGGTGGATCTTTGGCTTCGAGAGATCTTTTCATCTTTGACGAGGACTTCTGGGCGG 62
DQ 982 GGGGTGAACAGTAGGCTGGGAGAAATTCCTTCGACAAAGAGTTGGAGATGAT 1041
DB 63 GTATGTAACACTCTGGGCTCTGTGTG 100
DB 1042 GAATACAAACCCCTGGGACTGCTGCTGCTAGTG 1079

RESULT 9
US-09-764-870-609
; Sequence 609, Application US/09764870
; Patent No. US20020042386A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214
; CURRENT APPLICATION NUMBER: US/09/764,870
; CURRENT FILING DATE: 2001-01-17
; PRIOR APPLICATION DATA REMOVED - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 609
; LENGTH: 1314
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-870-609

Query Match          26.3%; Score 29.2; DB 10; Length 1314;
Best Local Similarity 56.1%; Pred. No. 1.6;
Matches 55; Conservative 0; Mismatches 43; Indels 0; Gaps 0;
```

NAME/KEY: SITE
LOCATION: 6286

; NAME/KEY: m

```

1 OTHER INFORMATION: n equals a,t,g, or c
2 NAME/KEY: misc feature
3 LOCATION: (340)
4 OTHER INFORMATION: n equals a,t,g, or c
5 NAME/KEY: misc feature
6 LOCATION: (370)
7 OTHER INFORMATION: n equals a,t,g, or c
8 NAME/KEY: misc feature
9 LOCATION: (359)
10 OTHER INFORMATION: n equals a,t,g, or c
11 NAME/KEY: misc feature
12 LOCATION: (402)
13 OTHER INFORMATION: n equals a,t,g, or c
14 NAME/KEY: misc feature
15 LOCATION: (403)
16 OTHER INFORMATION: n equals a,t,g, or c
17 NAME/KEY: misc feature
18 LOCATION: (406)
19 OTHER INFORMATION: n equals a,t,g, or c
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99
100
101
102
103
104
105
106
107
108
109
110
111
112
113
114
115
116
117
118
119
120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
166
167
168
169
170
171
172
173
174
175
176
177
178
179
180
181
182
183
184
185
186
187
188
189
190
191
192
193
194
195
196
197
198
199
200
201
202
203
204
205
206
207
208
209
210
211
212
213
214
215
216
217
218
219
220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240
241
242
243
244
245
246
247
248
249
250
251
252
253
254
255
256
257
258
259
260
261
262
263
264
265
266
267
268
269
270
271
272
273
274
275
276
277
278
279
280
281
282
283
284
285
286
287
288
289
290
291
292
293
294
295
296
297
298
299
300
301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340
341
342
343
344
345
346
347
348
349
350
351
352
353
354
355
356
357
358
359
360
361
362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445
446
447
448
449
450
451
452
453
454
455
456
457
458
459
460
461
462
463
464
465
466
467
468
469
470
471
472
473
474
475
476
477
478
479
480
481
482
483
484
485
486
487
488
489
490
491
492
493
494
495
496
497
498
499
500
501
502
503
504
505
506
507
508
509
510
511
512
513
514
515
516
517
518
519
520
521
522
523
524
525
526
527
528
529
530
531
532
533
534
535
536
537
538
539
540
541
542
543
544
545
546
547
548
549
550
551
552
553
554
555
556
557
558
559
560
561
562
563
564
565
566
567
568
569
570
571
572
573
574
575
576
577
578
579
580
581
582
583
584
585
586
587
588
589
590
591
592
593
594
595
596
597
598
599
600
601
602
603
604
605
606
607
608
609
610
611
612
613
614
615
616
617
618
619
620
621
622
623
624
625
626
627
628
629
630
631
632
633
634
635
636
637
638
639
640
641
642
643
644
645
646
647
648
649
650
651
652
653
654
655
656
657
658
659
660
661
662
663
664
665
666
667
668
669
670
671
672
673
674
675
676
677
678
679
680
681
682
683
684
685
686
687
688
689
690
691
692
693
694
695
696
697
698
699
700
701
702
703
704
705
706
707
708
709
710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727
728
729
730
731
732
733
734
735
736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807
808
809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828
829
830
831
832
833
834
835
836
837
838
839
840
841
842
843
844
845
846
847
848
849
850
851
852
853
854
855
856
857
858
859
860
861
862
863
864
865
866
867
868
869
870
871
872
873
874
875
876
877
878
879
880
881
882
883
884
885
886
887
888
889
890
891
892
893
894
895
896
897
898
899
900
901
902
903
904
905
906
907
908
909
910
911
912
913
914
915
916
917
918
919
920
921
922
923
924
925
926
927
928
929
930
931
932
933
934
935
936
937
938
939
940
941
942
943
944
945
946
947
948
949
950
951
952
953
954
955
956
957
958
959
960
961
962
963
964
965
966
967
968
969
970
971
972
973
974
975
976
977
978
979
980
981
982
983
984
985
986
987
988
989
990
991
992
993
994
995
996
997
998
999
1000
1001
1002
1003
1004
1005
1006
1007
1008
1009

```